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Kim et al.

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(54) **CONDENSED CYCLIC COMPOUND AND ORGANIC LIGHT-EMITTING DEVICE INCLUDING THE SAME**

(52) **U.S. Cl.**
CPC **H01L 51/0061** (2013.01); **C07D 221/18** (2013.01); **C07D 401/12** (2013.01);
(Continued)

(71) Applicant: **SAMSUNG DISPLAY CO., LTD.,**
Yongin-si (KR)

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See application file for complete search history.

(72) Inventors: **Sooyon Kim**, Yongin-si (KR);
Youngkook Kim, Yongin-si (KR);
Jongwoo Kim, Yongin-si (KR); **Jeonga Oh**, Yongin-si (KR); **Hyejin Jung**, Yongin-si (KR); **Seokhwan Hwang**, Yongin-si (KR); **Jaeyong Lee**, Yongin-si (KR)

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(73) Assignee: **Samsung Display Co., Ltd.**, Yongin-si (KR)

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(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 551 days.

This patent is subject to a terminal disclaimer.

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Primary Examiner — William E McClain

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(74) *Attorney, Agent, or Firm* — Lewis Roca Rothgerber Christie LLP

(30) **Foreign Application Priority Data**

Nov. 3, 2015 (KR) 10-2015-0153802

(57) **ABSTRACT**

An organic light-emitting device includes: a first electrode; a second electrode facing the first electrode; and an organic layer between the first electrode and the second electrode, the organic layer including an emission layer, wherein the organic layer includes a condensed cyclic compound of Formula 1:

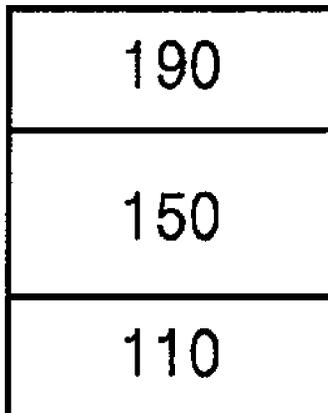
(51) **Int. Cl.**

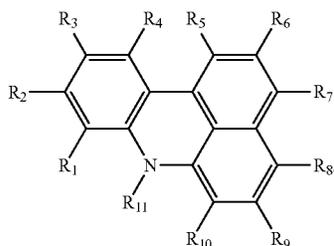
H01L 51/00 (2006.01)
C07D 405/14 (2006.01)

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Formula 1

An organic light-emitting device including the condensed cyclic compound of Formula 1 may have high efficiency, low driving voltage, high luminance, and a long lifespan.

13 Claims, 1 Drawing Sheet

(51) **Int. Cl.**

C07D 401/12 (2006.01)
C07D 221/18 (2006.01)
C07D 409/14 (2006.01)
H01L 51/50 (2006.01)

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CPC **C07D 405/14** (2013.01); **C07D 409/14** (2013.01); **H01L 51/006** (2013.01); **H01L 51/0052** (2013.01); **H01L 51/0072** (2013.01); **H01L 51/0073** (2013.01); **H01L 51/0074** (2013.01); **H01L 51/0056** (2013.01); **H01L 51/5056** (2013.01); **H01L 51/5088** (2013.01); **H01L 51/5096** (2013.01)

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150
110

**CONDENSED CYCLIC COMPOUND AND
ORGANIC LIGHT-EMITTING DEVICE
INCLUDING THE SAME**

CROSS-REFERENCE TO RELATED
APPLICATION

This application claims priority to and the benefit of Korean Patent Application No. 10-2015-0153802, filed on Nov. 3, 2015, in the Korean Intellectual Property Office, the entire content of which is incorporated herein by reference.

BACKGROUND

1. Field

One or more aspects of embodiments of the present disclosure relate to a compound for an organic light-emitting device, and an organic light-emitting device including the same.

2. Description of the Related Art

Organic light-emitting devices are self-emission devices that have wide viewing angles, high contrast ratios, short response times, and excellent brightness, driving voltage, and/or response speed characteristics, and can produce full-color images.

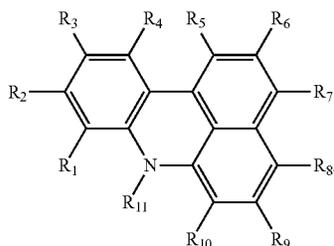
An organic light-emitting device may include a first electrode disposed (e.g., positioned) on a substrate, and a hole transport region, an emission layer, an electron transport region, and a second electrode, which are sequentially disposed on the first electrode. Holes provided from the first electrode, for example, may move toward the emission layer through the hole transport region, and electrons provided from the second electrode, for example, may move toward the emission layer through the electron transport region. Carriers, such as holes and electrons, can then recombine in the emission layer to produce excitons. These excitons transition from an excited state to a ground state, thereby generating light.

SUMMARY

One or more aspects of embodiments of the present disclosure are directed toward a condensed cyclic compound having a novel structure and an organic light-emitting device including the condensed cyclic compound.

Additional aspects will be set forth in part in the description which follows and, in part, will be apparent from the description, or may be learned by practice of the presented embodiments.

According to one or more embodiments, a condensed cyclic compound is represented by Formula 1:

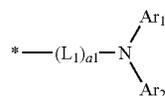


Formula 1

In Formula 1,

R_1 to R_{11} may each independently be selected from a group represented by Formula 2, hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a substituted or unsubstituted C_1 - C_{60} alkyl group, a substituted or unsubstituted C_2 - C_{60} alkenyl group, a substituted or unsubstituted C_2 - C_{60} alkenyl group, a substituted or unsubstituted C_1 - C_{60} alkoxy group, a substituted or unsubstituted C_3 - C_{10} cycloalkyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkyl group, a substituted or unsubstituted C_3 - C_{10} cycloalkenyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkenyl group, a substituted or unsubstituted C_6 - C_{60} aryl group, a substituted or unsubstituted C_6 - C_{60} aryloxy group, a substituted or unsubstituted C_6 - C_{60} arylthio group, a substituted or unsubstituted C_1 - C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, and —Si(Q_1)(Q_2)(Q_3),

provided that R_{11} is not the group represented by Formula 2, and a substituent selected from R_1 to R_{10} may be represented by Formula 2:



Formula 2

In Formula 2,

L_1 may be selected from a substituted or unsubstituted C_3 - C_{10} cycloalkylene group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkylene group, a substituted or unsubstituted C_3 - C_{10} cycloalkenylene group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkenylene group, a substituted or unsubstituted C_6 - C_{60} arylene group, a substituted or unsubstituted C_1 - C_{60} heteroarylene group, a substituted or unsubstituted divalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted divalent non-aromatic condensed heteropolycyclic group;

a_1 may be an integer selected from 0 to 3, wherein when a_1 is two or more, two or more L_1 (s) may be identical to or different from each other;

Ar_1 and Ar_2 may each independently be selected from a substituted or unsubstituted C_3 - C_{10} cycloalkyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkyl group, a substituted or unsubstituted C_3 - C_{10} cycloalkenyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkenyl group, a substituted or unsubstituted C_6 - C_{60} aryl group, a substituted or unsubstituted C_1 - C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, provided that at least one selected from Ar_1 and Ar_2 may each independently be selected from a substituted or unsubstituted fluorenyl group, a substituted or unsubstituted spiro-bifluorenyl group, a substituted or unsubstituted carbazolyl group, a substituted or unsubstituted dibenzofuranyl group, a substituted or unsubstituted dibenzothiophenyl group, and a substituted or unsubstituted dibenzosilolyl group.

At least one substituent of the substituted fluorenyl group, substituted spiro-bifluorenyl group, a substituted carbazolyl group, a substituted dibenzofuranyl group, a substituted

dibenzothiophenyl group, a substituted dibenzosilolyl group, substituted C₃-C₁₀ cycloalkylene group, substituted C₁-C₁₀ heterocycloalkylene group, substituted C₃-C₁₀ cycloalkenylene group, substituted C₁-C₁₀ heterocycloalkenylene group, substituted C₆-C₆₀ arylene group, substituted C₁-C₆₀ heteroarylene group, a substituted divalent non-aromatic condensed polycyclic group, a substituted divalent non-aromatic condensed heteropolycyclic group, substituted C₁-C₆₀ alkyl group, substituted C₂-C₆₀ alkenyl group, substituted C₂-C₆₀ alkynyl group, substituted C₁-C₆₀ alkoxy group, substituted C₃-C₁₀ cycloalkyl group, substituted C₁-C₁₀ heterocycloalkyl group, substituted C₃-C₁₀ cycloalkenyl group, substituted C₁-C₁₀ heterocycloalkenyl group, substituted C₆-C₆₀ aryl group, substituted C₆-C₆₀ aryloxy group, substituted C₆-C₆₀ arylthio group, substituted C₁-C₆₀ heteroaryl group, substituted monovalent non-aromatic condensed polycyclic group, and substituted monovalent non-aromatic condensed heteropolycyclic group may be selected from the group consisting of:

deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group,

a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, C₆-C₆₀ arylthio group, C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, and —Si(Q₁₁)(Q₁₂)(Q₁₃),

a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, a biphenyl group, and a terphenyl group,

a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, a biphenyl group, and a terphenyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a

monovalent non-aromatic condensed heteropolycyclic group, a biphenyl group, a terphenyl group, and —Si(Q₂₁)(Q₂₂)(Q₂₃), and

—Si(Q₃₁)(Q₃₂)(Q₃₃),
 wherein Q₁ to Q₃, Q₁₁ to Q₁₃, Q₂₁ to Q₂₃, and Q₃₁ to Q₃₃ may each independently be selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, a biphenyl group, and a terphenyl group.

According to one or more embodiments, an organic light-emitting device includes a first electrode; a second electrode facing the first electrode; and an organic layer between the first electrode and the second electrode, the organic layer including an emission layer, wherein the organic layer includes the condensed cyclic compound as described above.

BRIEF DESCRIPTION OF THE DRAWING

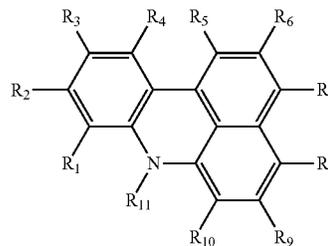
These and/or other aspects will become apparent and more readily appreciated from the following description of the embodiments, taken in conjunction with the drawing, which is a schematic diagram of a structure of an organic light-emitting device according to an embodiment of the present disclosure.

DETAILED DESCRIPTION

Reference will now be made in more detail to embodiments, examples of which are illustrated in the accompanying drawing, wherein like reference numerals refer to like elements throughout. In this regard, the present embodiments may have different forms and should not be construed as being limited to the descriptions set forth herein. Accordingly, the embodiments are merely described below, by referring to the drawing, to explain aspects of the present description. As used herein, the term “and/or” includes any and all combinations of one or more of the associated listed items. Expressions such as “at least one of,” “one of,” and “selected from,” when preceding a list of elements, modify the entire list of elements and do not modify the individual elements of the list. Further, the use of “may” when describing embodiments of the present invention refers to “one or more embodiments of the present invention.”

A condensed cyclic compound according to embodiments of the present disclosure is represented by Formula 1:

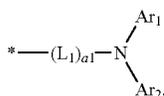
Formula 1



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R₁ to R₁₁ in Formula 1 may each independently be selected from a group represented by Formula 2, hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a substituted or unsubstituted C₁-C₆₀ alkyl group, a substituted or unsubstituted C₂-C₆₀ alkenyl group, a substituted or unsubstituted C₂-C₆₀ alkynyl group, a substituted or unsubstituted C₁-C₆₀ alkoxy group, a substituted or unsubstituted C₃-C₁₀ cycloalkyl group, a substituted or unsubstituted C₁-C₁₀ heterocycloalkyl group, a substituted or unsubstituted C₃-C₁₀ cycloalkenyl group, a substituted or unsubstituted C₁-C₁₀ heterocycloalkenyl group, a substituted or unsubstituted C₆-C₆₀ aryl group, a substituted or unsubstituted C₆-C₆₀ aryloxy group, a substituted or unsubstituted C₆-C₆₀ arylthio group, a substituted or unsubstituted C₁-C₆₀ heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, and —Si(Q₁)(Q₂)(Q₃), provided that R₁₁ is not the group represented by Formula 2.

A substituent selected from R₁ to R₁₀ in Formula 1 may be the group represented by Formula 2.



Formula 2

In Formula 2,

L₁ may be selected from a substituted or unsubstituted C₃-C₁₀ cycloalkylene group, a substituted or unsubstituted C₁-C₁₀ heterocycloalkylene group, a substituted or unsubstituted C₃-C₁₀ cycloalkenylene group, a substituted or unsubstituted C₁-C₁₀ heterocycloalkenylene group, a substituted or unsubstituted C₆-C₆₀ arylene group, a substituted or unsubstituted C₁-C₆₀ heteroarylene group, a substituted or unsubstituted divalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted divalent non-aromatic condensed heteropolycyclic group.

For example, L₁ in Formula 2 may be selected from the group consisting of:

a phenylene group, a pentalenylene group, an indenylene group, a naphthylene group, an azulenylenylene group, a heptalenylene group, an indacenylenylene group, an acenaphthylene group, a fluorenylenylene group, spiro-bifluorenylenylene group, a benzofluorenylenylene group, a dibenzofluorenylenylene group, a phenalenylene group, a phenanthrenylene group, an anthracenylenylene group, a fluoranthrenylene group, a triphenylenylene group, a pyrenylene group, a chrysenylene group, a naphthacenylenylene group, a picenylene group, a perylenylene group, a pentaphenylenylene group, a hexacenylenylene group, a pentacenylenylene group, a rubicenylenylene group, a coronenylenylene group, an ovalenylenylene group, a pyrrolylenylene group, a thiophenylenylene group, a furanylenylene group, an imidazolylene group, a pyrazolylene group, a thiazolylene group, an isothiazolylene group, an oxazolylene group, an isoxazolylene group, a pyridinylenylene group, a pyrazinylenylene group, a pyrimidinylene group, a pyridazinylenylene group, an isoindolylenylene group, an indolylenylene group, an indazolylene group, a purinylenylene group, a quinolinylene group, an isoquinolinylene group, a benzoquinolinylene group, a phthalazinylene group, a naphthyridinylene group, a quinoxalinylene group, a quinazolinylene group, a cinnolinylene group, a carba-

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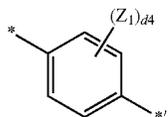
zolylene group, a phenanthridinylene group, an acridinylene group, a phenanthrolinylene group, a phenazinylene group, a benzoimidazolylene group, a benzofuranylenylene group, a benzothioiophenylene group, an isobenzothiazolylene group, a benzoxazolylene group, an isobenzoxazolylene group, a triazolylene group, a tetrazolylene group, an oxadiazolylene group, a triazinylene group, a dibenzofuranylenylene group, a dibenzothiophenylene group, a dibenzosilolylene group, a benzocarbazolylene group, a dibenzocarbazolylene group, a thiadiazolylene group, an imidazopyridinylene group, and an imidazopyrimidinylene group, and

a phenylene group, a pentalenylene group, an indenylene group, a naphthylene group, an azulenylenylene group, a heptalenylene group, an indacenylenylene group, an acenaphthylene group, a fluorenylenylene group, spiro-bifluorenylenylene group, a benzofluorenylenylene group, a dibenzofluorenylenylene group, a phenalenylene group, a phenanthrenylene group, an anthracenylenylene group, a fluoranthrenylene group, a triphenylenylene group, a pyrenylene group, a chrysenylene group, a naphthacenylenylene group, a picenylene group, a perylenylene group, a pentaphenylenylene group, a hexacenylenylene group, a pentacenylenylene group, a rubicenylenylene group, a coronenylenylene group, an ovalenylenylene group, a pyrrolylenylene group, a thiophenylenylene group, a furanylenylene group, an imidazolylene group, a pyrazolylene group, a thiazolylene group, an isothiazolylene group, an oxazolylene group, an isoxazolylene group, a pyridinylenylene group, a pyrazinylenylene group, a pyrimidinylene group, a pyridazinylenylene group, an isoindolylenylene group, an indolylenylene group, an indazolylene group, a purinylenylene group, a quinolinylene group, an isoquinolinylene group, a benzoquinolinylene group, a phthalazinylene group, a naphthyridinylene group, a quinoxalinylene group, a quinazolinylene group, a cinnolinylene group, a carbazolylene group, a phenanthridinylene group, an acridinylene group, a phenanthrolinylene group, a phenazinylene group, a benzoimidazolylene group, a benzofuranylenylene group, a benzothioiophenylene group, an isobenzothiazolylene group, a benzoxazolylene group, an isobenzoxazolylene group, a triazolylene group, a tetrazolylene group, an oxadiazolylene group, a triazinylene group, a dibenzofuranylenylene group, a dibenzothiophenylene group, a dibenzosilolylene group, a benzocarbazolylene group, a dibenzocarbazolylene group, a thiadiazolylene group, an imidazopyridinylene group, and an imidazopyrimidinylene group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a cyclopentyl group, a cyclohexyl group, a cycloheptyl group, a cyclopentenyl group, a cyclohexenyl group, a phenyl group, a biphenyl group, a terphenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an azulenyl group, a heptalenyl group, an indacenyl group, an acenaphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenalenyl group, a phenanthrenyl group, an anthracenyl group, a fluoranthrenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a naphthacenyl group, a picenyl group, a perylenyl group, a pentaphenyl group, a hexacenyl group, a pentacenyl group, a rubicenyl group, a coronenyl group, an ovalenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group,

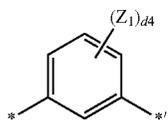
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an indolyl group, an indazolyl group, a purinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a phthalazinyl group, a naphthyridinyl group, a quinoxalinyl group, a quinazolinyl group, a cinnolinyl group, a carbazolyl group, a phenanthridinyl group, an acridinyl group, a phenanthrolinyl group, a phenazinyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolylene group, a benzocarbazolyl group, a dibenzocarbazolyl group, a thiadiazolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group.

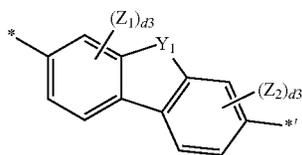
In various embodiments, L_1 in Formula 2 may be selected from groups represented by Formula 3-1 to Formula 3-42:



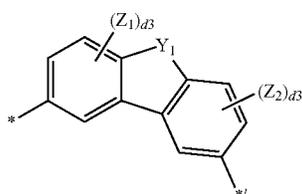
Formula 3-1



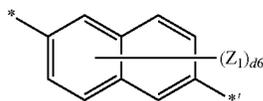
Formula 3-2



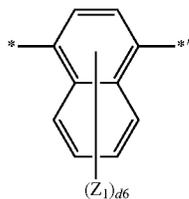
Formula 3-3



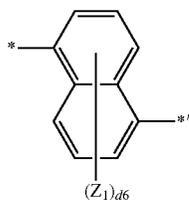
Formula 3-4



Formula 3-5



Formula 3-6

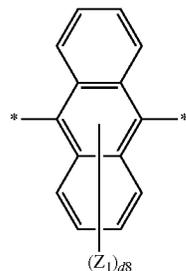


Formula 3-7

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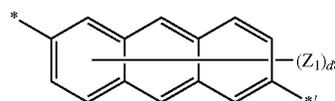
Formula 3-8



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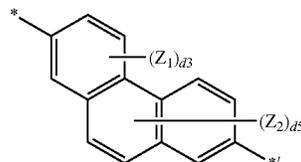
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Formula 3-9



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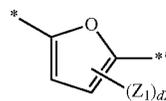
Formula 3-10



Formula 3-1

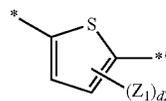
Formula 3-2

Formula 3-11



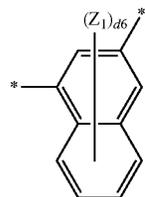
Formula 3-3

Formula 3-12



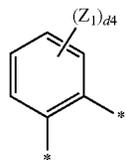
Formula 3-4

Formula 3-13



Formula 3-5

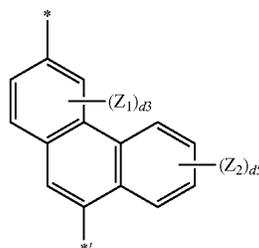
Formula 3-14



Formula 3-6

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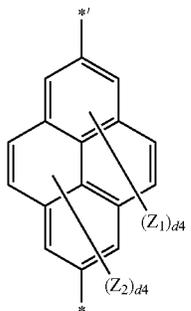
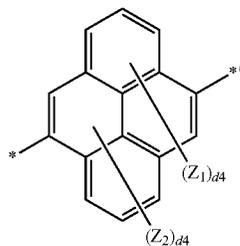
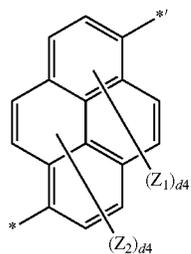
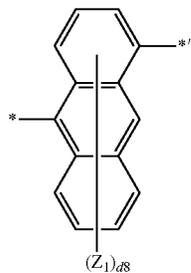
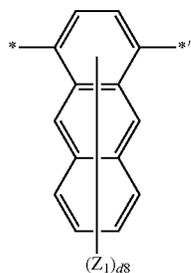
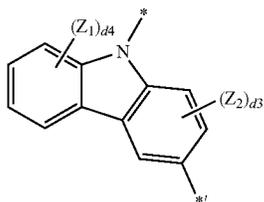
Formula 3-15



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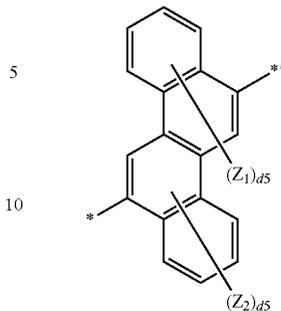
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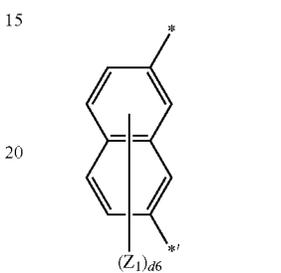


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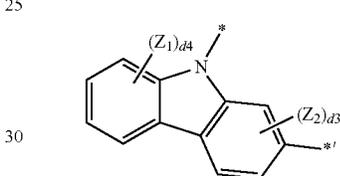
Formula 3-16



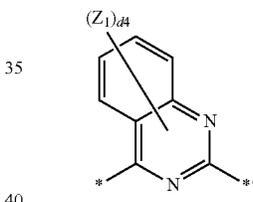
Formula 3-17



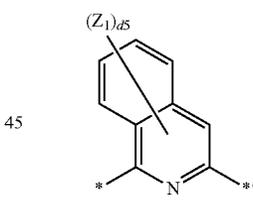
Formula 3-18



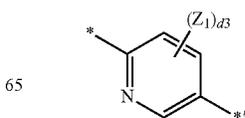
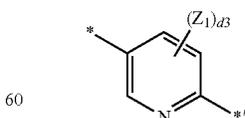
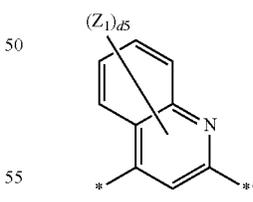
Formula 3-19



Formula 3-20



Formula 3-21



Formula 3-22

Formula 3-23

Formula 3-24

Formula 3-25

Formula 3-26

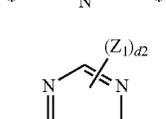
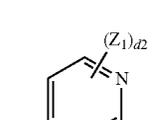
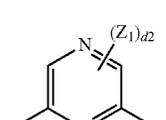
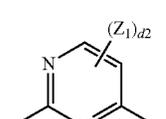
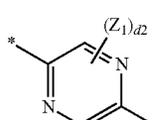
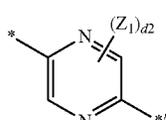
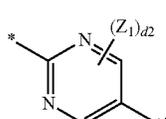
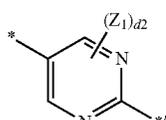
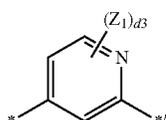
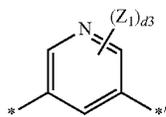
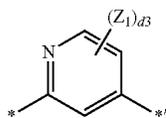
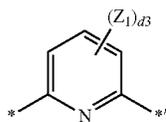
Formula 3-27

Formula 3-28

Formula 3-29

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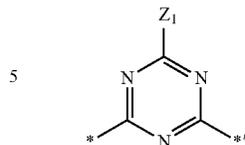
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12

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Formula 3-30



Formula 3-31

10 In Formulae 3-1 to 3-42,

Y_1 may be selected from O, S, $C(Z_3)(Z_4)$, $N(Z_5)$, and $Si(Z_6)(Z_7)$;

Formula 3-32

Z_1 to Z_7 may each independently be selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{20} alkyl group, a C_1 - C_{20} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazoliny group, a carbazolyl group, a triazinyl group, and — $Si(Q_{31})(Q_{32})(Q_{33})$;

Formula 3-33

Formula 3-34

Formula 3-35

Q_{31} to Q_{33} may each independently be selected from a C_1 - C_{10} alkyl group, a C_1 - C_{10} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, and a naphthyl group;

Formula 3-36

d_2 may be 1 or 2;
 d_3 may be an integer selected from 1 to 3;
 d_4 may be an integer selected from 1 to 4;
 d_5 may be an integer selected from 1 to 5;
 d_6 may be an integer selected from 1 to 6;
 d_8 may be an integer selected from 1 to 8; and
 * and *' each indicate a binding site to a neighboring atom.

Formula 3-37

In various embodiments, L_1 in Formula 2 may be selected from the group consisting of:

Formula 3-38

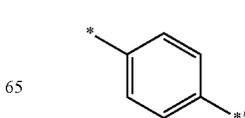
a phenylene group, a naphthylene group, a fluorenylene group, a phenanthrenylene group, an anthracenylylene group, a triphenylylene group, and a pyrenylene group, and
 a phenylene group, a naphthylene group, a fluorenylene group, a phenanthrenylene group, an anthracenylylene group, a triphenylylene group, and a pyrenylene group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{20} alkyl group, a C_1 - C_{20} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, and a naphthyl group, but L_1 is not limited thereto.

Formula 3-39

Formula 3-40

In various embodiments, L_1 in Formula 2 may be selected from groups represented by Formula 4-1 to Formula 4-29, but is not limited thereto:

Formula 3-41

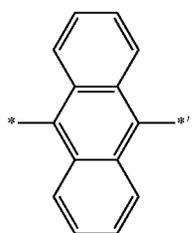
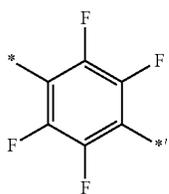
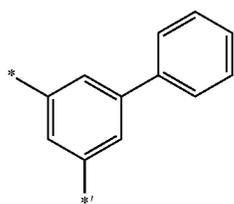
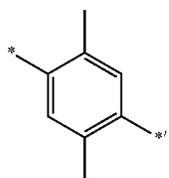
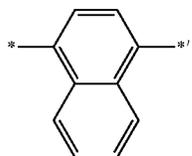
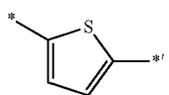
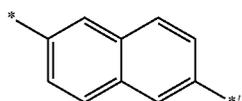
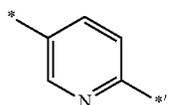
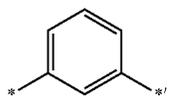
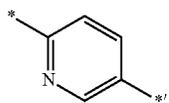


Formula 3-42

Formula 4-1

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13
-continued



14
-continued

Formula 4-2

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Formula 4-3

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Formula 4-4

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Formula 4-5

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Formula 4-6

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Formula 4-7

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Formula 4-8

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Formula 4-9

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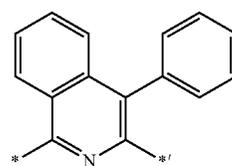
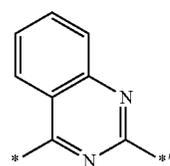
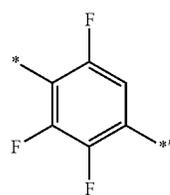
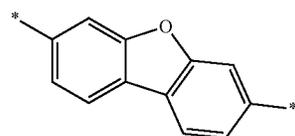
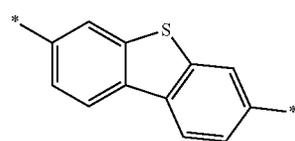
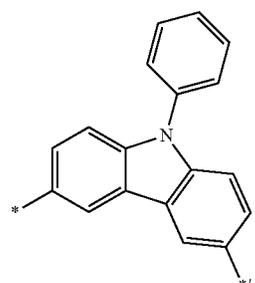
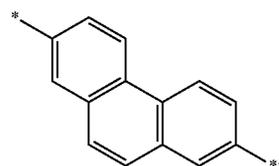
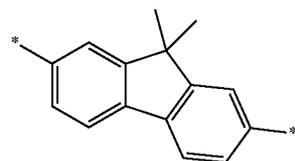
Formula 4-10

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Formula 4-11

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Formula 4-12

Formula 4-13

Formula 4-14

Formula 4-15

Formula 4-16

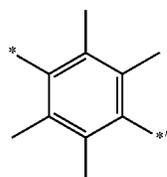
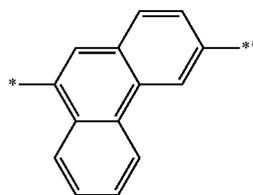
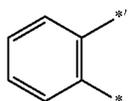
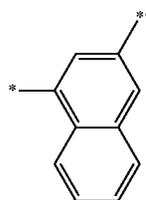
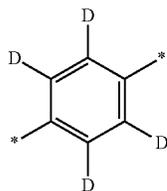
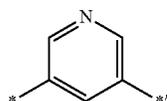
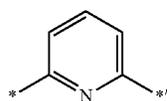
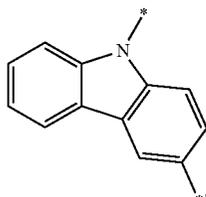
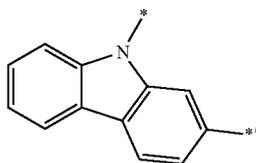
Formula 4-17

Formula 4-18

Formula 4-19

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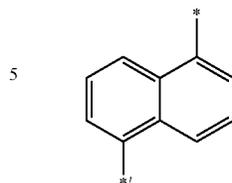
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16

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Formula 4-20



Formula 4-21 10

* and *' in Formulae 4-1 to 4-29 each indicate a binding site to a neighboring atom, and "D" may refer to deuterium.

Formula 4-22 15

a1 in Formula 2 may be an integer selected from 0 to 3. a1 indicates the number of L₁(s) in Formula 2, and when a1 is two or more, two or more L₁(s) may be identical to or different from each other. For example, when a1 is 0, *-(L₁)_{a1}-*' indicates a single bond. In various embodiments, a1 may be 0, 1, or 2. For example, a1 may be 0 or 1.

Formula 4-23 20

Ar₁ and Ar₂ in Formula 2 may each independently be selected from a substituted or unsubstituted C₃-C₁₀ cycloalkyl group, a substituted or unsubstituted C₁-C₁₀ heterocycloalkyl group, a substituted or unsubstituted C₃-C₁₀ cycloalkenyl group, a substituted or unsubstituted C₁-C₁₀ heterocycloalkenyl group, a substituted or unsubstituted C₆-C₆₀ aryl group, a substituted or unsubstituted C₁-C₆₀ heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, provided that at least one selected from Ar₁ and Ar₂ may each independently be selected from a substituted or unsubstituted fluorenyl group, a substituted or unsubstituted spiro-bifluorenyl group, a substituted or unsubstituted carbazolyl group, a substituted or unsubstituted dibenzofuranyl group, a substituted or unsubstituted dibenzothiophenyl group, and a substituted or unsubstituted dibenzosilolyl group.

Formula 4-24 25

Formula 4-25 30

Formula 4-26 35

Formula 4-27 40

Formula 4-28 45

Formula 4-29 50

For example, Ar₁ and Ar₂ in Formula 2 may each independently be selected from:

a substituted or unsubstituted phenyl group, a substituted or unsubstituted biphenyl group, a substituted or unsubstituted terphenyl group, a substituted or unsubstituted pentalenyl group, a substituted or unsubstituted indenyl group, a substituted or unsubstituted naphthyl group, a substituted or unsubstituted azulenyl group, a substituted or unsubstituted heptalenyl group, a substituted or unsubstituted indacenyl group, a substituted or unsubstituted acenaphthyl group, a substituted or unsubstituted fluorenyl group, a substituted or unsubstituted spiro-bifluorenyl group, a substituted or unsubstituted benzofluorenyl group, a substituted or unsubstituted dibenzofluorenyl group, a substituted or unsubstituted phenalenyl group, a substituted or unsubstituted phenanthrenyl group, a substituted or unsubstituted anthracenyl group, a substituted or unsubstituted fluoranthenyl group, a substituted or unsubstituted triphenylenyl group, a substituted or unsubstituted pyrenyl group, a substituted or unsubstituted chrysenyl group, a substituted or unsubstituted naphthacenyl group, a substituted or unsubstituted picenyl group, a substituted or unsubstituted perylenyl group, a substituted or unsubstituted pentaphenyl group, a substituted or unsubstituted hexacenyl group, a substituted or unsubstituted pentacenyl group, a substituted or unsubstituted rubicenyl group, a substituted or unsubstituted coronenyl group, a substituted or unsubstituted ovalenyl group, a substituted or unsubstituted pyrrolyl group, a substituted or unsubstituted thiophenyl group, a substituted or unsubstituted furanyl group, a substituted or unsubstituted imidazolyl group, a

substituted or unsubstituted pyrazolyl group, a substituted or unsubstituted thiazolyl group, a substituted or unsubstituted isothiazolyl group, a substituted or unsubstituted oxazolyl group, a substituted or unsubstituted isoxazolyl group, a substituted or unsubstituted pyridinyl group, a substituted or unsubstituted pyrazinyl group, a substituted or unsubstituted pyrimidinyl group, a substituted or unsubstituted pyridazinyl group, a substituted or unsubstituted isoindolyl group, a substituted or unsubstituted indolyl group, a substituted or unsubstituted indazolyl group, a substituted or unsubstituted purinyl group, a substituted or unsubstituted quinolinyl group, a substituted or unsubstituted isoquinolinyl group, a substituted or unsubstituted benzoquinolinyl group, a substituted or unsubstituted phthalazinyl group, a substituted or unsubstituted naphthyridinyl group, a substituted or unsubstituted quinoxalinyl group, a substituted or unsubstituted quinazoliny group, a substituted or unsubstituted cinnolinyl group, a substituted or unsubstituted carbazolyl group, a substituted or unsubstituted phenanthridinyl group, a substituted or unsubstituted acridinyl group, a substituted or unsubstituted phenanthrolinyl group, a substituted or unsubstituted phenaziny group, a substituted or unsubstituted benzoimidazolyl group, a substituted or unsubstituted benzofuranyl group, a substituted or unsubstituted dibenzofuranyl group, a substituted or unsubstituted benzothiophenyl group, a substituted or unsubstituted dibenzothiophenyl group, a substituted or unsubstituted isobenzothiazolyl group, a substituted or unsubstituted benzoxazolyl group, a substituted or unsubstituted isobenzoxazolyl group, a substituted or unsubstituted triazolyl group, a substituted or unsubstituted tetrazolyl group, a substituted or unsubstituted oxadiazolyl group, a substituted or unsubstituted triazinyl group, a substituted or unsubstituted benzocarbazolyl group, a substituted or unsubstituted dibenzocarbazolyl group, the substituted thiadiazolyl group, the substituted imidazopyridinyl group, and a substituted or unsubstituted imidazopyrimidinyl group,

where at least one substituent of the substituted phenyl group, the substituted biphenyl group, the substituted terphenyl group, the substituted pentalenyl group, the substituted indenyl group, the substituted naphthyl group, the substituted azulenyl group, the substituted heptalenyl group, the substituted indacenyl group, the substituted acenaphthyl group, the substituted fluorenyl group, substituted spiro-bifluorenyl group, the substituted benzofluorenyl group, the substituted dibenzofluorenyl group, the substituted phenalenyl group, the substituted phenanthrenyl group, the substituted anthracenyl group, the substituted fluoranthenyl group, the substituted triphenylenyl group, the substituted pyrenyl group, the substituted chrysenyl group, the substituted naphthacenyl group, the substituted picenyl group, the substituted perylenyl group, the substituted pentaphenyl group, the substituted hexacenyl group, the substituted pentacenyl group, the substituted rubicenyl group, the substituted coronenyl group, the substituted ovalenyl group, the substituted pyrrolyl group, the substituted thiophenyl group, the substituted furanyl group, the substituted imidazolyl group, the substituted pyrazolyl group, the substituted thiazolyl group, the substituted isothiazolyl group, the substituted oxazolyl group, the substituted isoxazolyl group, the substituted pyridinyl group, the substituted pyrazinyl group, the substituted pyrimidinyl group, the substituted pyridazinyl group, the substituted isoindolyl group, the substituted indolyl group, the substituted indazolyl group, the substituted purinyl group, a substituted quinolinyl group, the substituted isoquinolinyl group, the substituted benzoquinolinyl group, the substituted phthalazinyl group, the substituted naphthy-

ridinyl group, the substituted quinoxalinyl group, the substituted quinazoliny group, the substituted cinnolinyl group, the substituted carbazolyl group, the substituted phenanthridinyl group, the substituted acridinyl group, the substituted phenanthrolinyl group, the substituted phenaziny group, the substituted benzoimidazolyl group, the substituted benzofuranyl group, the substituted dibenzofuranyl group, the substituted benzothiophenyl group, the substituted dibenzothiophenyl group, the substituted isobenzothiazolyl group, the substituted benzoxazolyl group, the substituted isobenzoxazolyl group, the substituted triazolyl group, the substituted tetrazolyl group, the substituted oxadiazolyl group, the substituted triazinyl group, the substituted benzocarbazolyl group, the substituted dibenzocarbazolyl group, the substituted thiadiazolyl group, the substituted imidazopyridinyl group, and the substituted imidazopyrimidinyl group may be selected from the group consisting of:

deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, and a C₁-C₂₀ alkoxy group,

a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, and a phosphoric acid group or a salt thereof,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a carbazolyl group, a benzocarbazolyl group, and a dibenzocarbazolyl group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a pyridinyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a carbazolyl group, a benzocarbazolyl group and a dibenzocarbazolyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group and a pyridinyl group, and

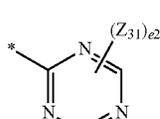
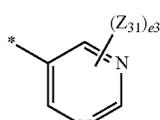
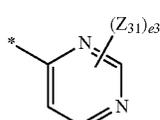
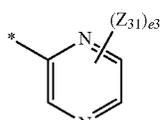
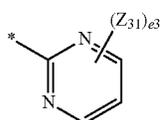
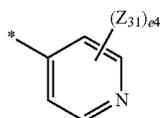
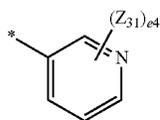
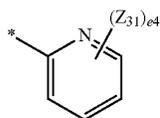
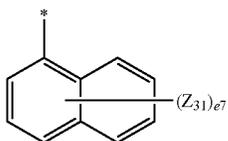
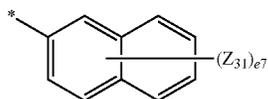
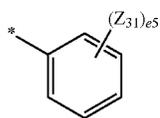
—Si(Q₃₁)(Q₃₂)(Q₃₃),

wherein Q₃₁ to Q₃₃ may each independently be selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, and a carbazolyl group.

In various embodiments, Ar₁ and Ar₂ in Formula 2 may each independently be selected from groups represented by Formula 5-1 to Formula 5-18, provided that at least one

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selected from Ar₁ and Ar₂ may each independently be selected from groups represented by Formulae 5-14 to 5-18:

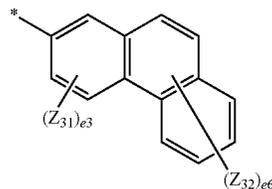


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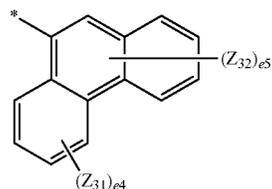
-continued

Formula 5-12

Formula 5-1 5



Formula 5-2 10



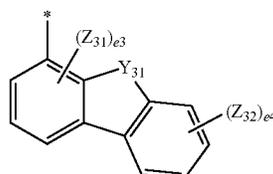
Formula 5-13

Formula 5-3 15

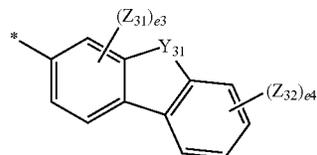


Formula 5-14

Formula 5-4 20

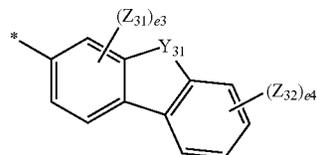


Formula 5-5 25



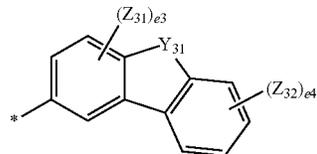
Formula 5-15

Formula 5-6 30



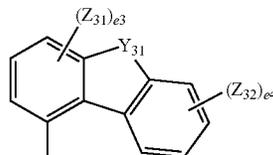
Formula 5-16

Formula 5-7 35



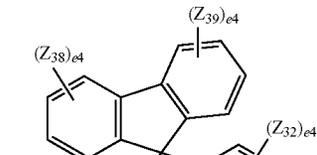
Formula 5-17

Formula 5-8 40

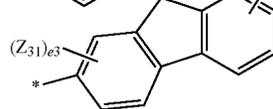


Formula 5-18

Formula 5-9 45



Formula 5-10 50



Formula 5-11 55

In Formulae 5-1 to 5-18,
 Y₃₁ may be selected from O, S, C(Z₃₃)(Z₃₄), N(Z₃₅), and Si(Z₃₆)(Z₃₇);

Z₃₁ to Z₃₉ may each independently be selected from the group consisting of:

hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group

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or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, and a C₁-C₂₀ alkoxy group,

a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, and a phosphoric acid group or a salt thereof,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a carbazolyl group, a benzocarbazolyl group and a dibenzocarbazolyl group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a pyridinyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a carbazolyl group, a benzocarbazolyl group, and a dibenzocarbazolyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, and a pyridinyl group, and

—Si(Q₃₁)(Q₃₂)(Q₃₃),

wherein Q₃₁ to Q₃₃ may each independently be selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, and a carbazolyl group;

e2 may be 1 or 2;

e3 may be an integer selected from 1 to 3;

e4 may be an integer selected from 1 to 4;

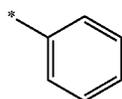
e5 may be an integer selected from 1 to 5;

e6 may be an integer selected from 1 to 6;

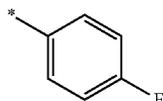
e7 may be an integer selected from 1 to 7; and

* indicates a binding site to a neighboring atom.

In various embodiments, Ar₁ and Ar₂ in Formula 2 may each independently be selected from groups represented by Formula 6-1 to Formula 6-42, provided that at least one selected from Ar₁ and Ar₂ may each independently be selected from groups represented by Formula 6-25 to Formula 6-42:



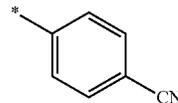
Formula 6-1



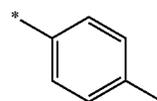
Formula 6-2

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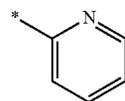
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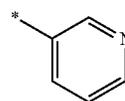
Formula 6-3



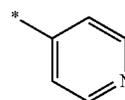
Formula 6-4



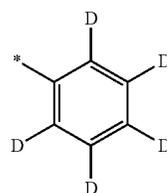
Formula 6-5



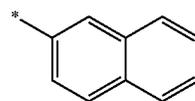
Formula 6-6



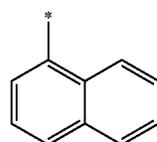
Formula 6-7



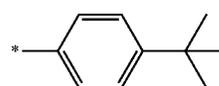
Formula 6-8



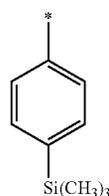
Formula 6-9



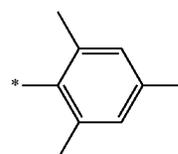
Formula 6-10



Formula 6-11

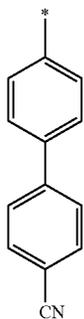
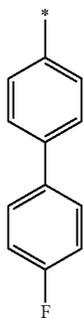
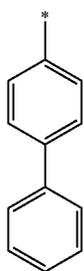
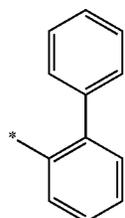
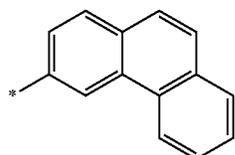
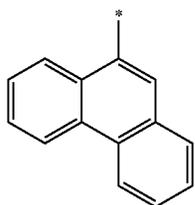


Formula 6-12



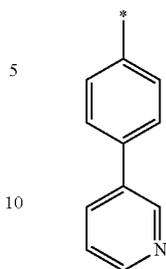
Formula 6-13

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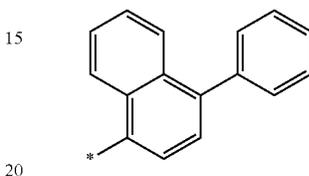


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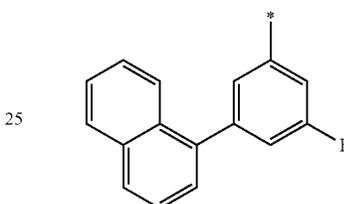
Formula 6-14



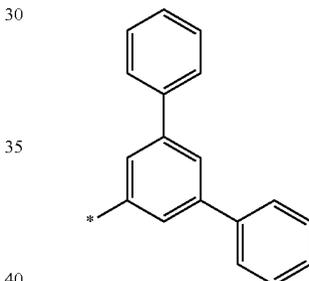
Formula 6-15



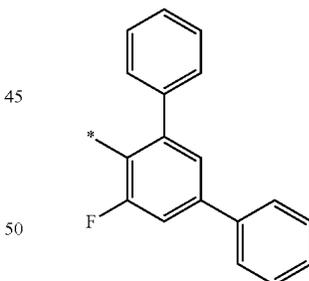
Formula 6-16



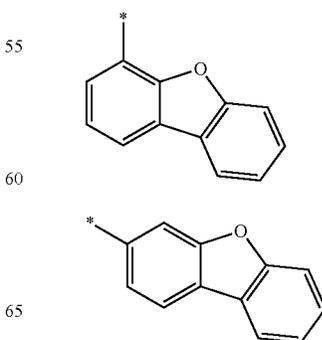
Formula 6-17



Formula 6-18



Formula 6-19



Formula 6-20

Formula 6-21

Formula 6-22

Formula 6-23

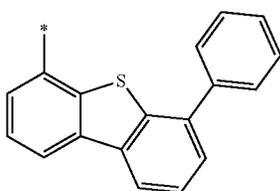
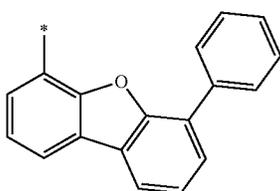
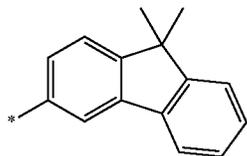
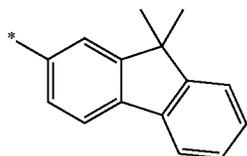
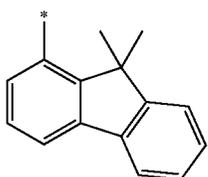
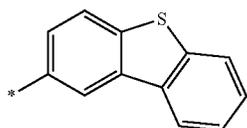
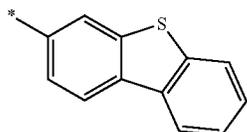
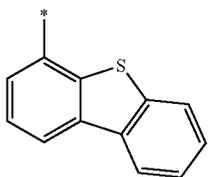
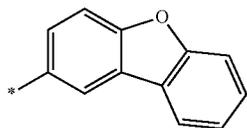
Formula 6-24

Formula 6-25

Formula 6-26

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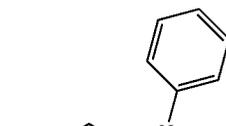


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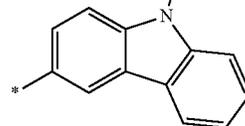
Formula 6-27

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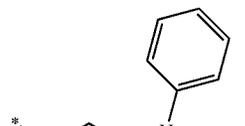
Formula 6-28

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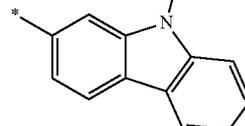


Formula 6-29

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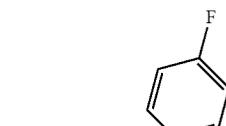


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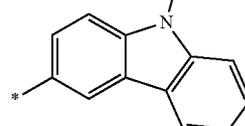
Formula 6-30

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Formula 6-31

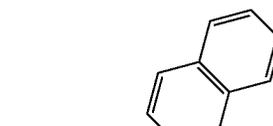
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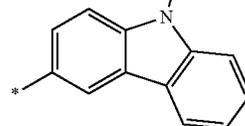
Formula 6-32

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Formula 6-33

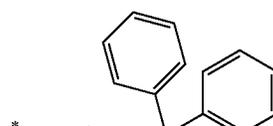
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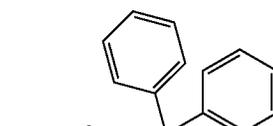
Formula 6-34

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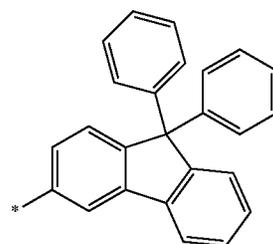


Formula 6-35

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Formula 6-36

Formula 6-37

Formula 6-38

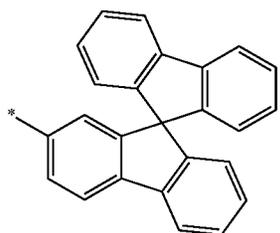
Formula 6-39

Formula 6-40

Formula 6-41

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* in Formulae 6-1 to 6-42 indicates a binding site to a neighboring atom, and "D" may refer to deuterium.

R₁ to R₁₀ in Formula 1 may each independently be selected from the group consisting of:

the group represented by Formula 2, hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, and a C₁-C₂₀ alkoxy group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₁₀ alkyl group, a C₁-C₁₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluore-

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Formula 6-42

nyl group, a phenanthrenyl group, an anthracenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, an imidazopyridinyl group, an imidazopyrimidinyl group, and —Si(Q₃₁)(Q₃₂)(Q₃₃), and

—Si(Q₁)(Q₂)(Q₃),

wherein Q₁ to Q₃ and Q₃₁ to Q₃₃ may each independently be selected from a C₁-C₁₀ alkyl group, a C₁-C₁₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, and a naphthyl group, but R₁ to R₁₀ are not limited thereto.

In various embodiments, R₁ to R₁₀ may each independently be selected from the group consisting of:

the group represented by Formula 2, hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₁₀ alkyl group, and a C₁-C₁₀ alkoxy group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a pyridinyl group, a pyrimidinyl group, and a triazinyl group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a pyridinyl group, a pyrimidinyl group, and a triazinyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₁₀ alkyl group, a C₁-C₁₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a pyridinyl group, a pyrimidinyl group, a triazinyl group, and —Si(Q₃₁)(Q₃₂)(Q₃₃), and

—Si(Q₁)(Q₂)(Q₃),

wherein Q₁ to Q₃ and Q₃₁ to Q₃₃ may each independently be selected from a C₁-C₁₀ alkyl group, a C₁-C₁₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, and a naphthyl group, but R₁ to R₁₀ are not limited thereto.

In various embodiments, R₁₁ may be selected from the group consisting of:

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl

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group, a dibenzothiophenyl group, a dibenzosilolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group, and

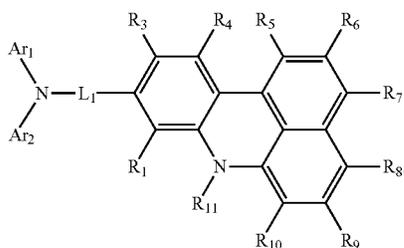
a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₁₀ alkyl group, a C₁-C₁₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, an imidazopyridinyl group, an imidazopyrimidinyl group, and —Si(Q₃₁)(Q₃₂)(Q₃₃), where Q₃₁ to Q₃₃ are as defined above, but R₁₁ is not limited thereto.

A substituent selected from R₁ to R₁₀ in Formula 1 may be the group represented by Formula 2.

In various embodiments, R₂ or R₇ in Formula 1 may be the group represented by Formula 2.

In various embodiments, the condensed cyclic compound represented by Formula 1 may be represented by one of Formulae 1-1(A), 1-1(B), 1-2(A), and 1-2(B), but is not limited thereto:

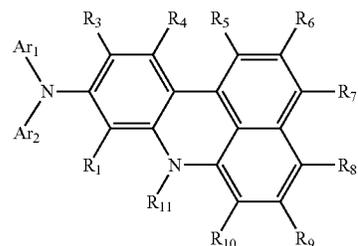
Formula 1-1(A)



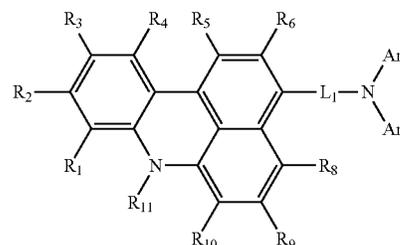
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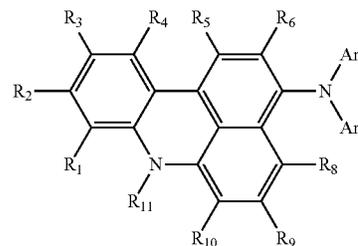
Formula 1-1(B)



Formula 1-2(A)



Formula 1-2(B)



L₁, Ar₁, Ar₂, and R₁ to R₁₁ in the Formulae 1-1(A), 1-1(B), 1-2(A), and 1-2(B) are the same as described above.

In various embodiments,

in the Formulae 1-1(A), 1-1(B), 1-2(A), and 1-2(B) above,

R₁ to R₁₀ may each be hydrogen;

L₁ may be selected from a substituted or unsubstituted phenylene group, a substituted or unsubstituted naphthylene group, a substituted or unsubstituted fluorenylene group, a substituted or unsubstituted phenanthrenylene group, a substituted or unsubstituted anthracenylene group, a substituted or unsubstituted triphenylenylene group, and a substituted or unsubstituted pyrenylene group;

R₁₁, Ar₁, and Ar₂ may each independently be selected from a substituted or unsubstituted phenyl group, a substituted or unsubstituted biphenyl group, a substituted or unsubstituted terphenyl group, a substituted or unsubstituted naphthyl group, a substituted or unsubstituted fluorenyl group, a substituted or unsubstituted phenanthrenyl group, a substituted or unsubstituted anthracenyl group, a substituted or unsubstituted triphenylenyl group, a substituted or unsubstituted pyrenyl group, a substituted or unsubstituted fluorenyl group, a substituted or unsubstituted spiro-bifluorenyl group, a substituted or unsubstituted carbazolyl group, a substituted or unsubstituted dibenzofuranyl group, a substituted or unsubstituted dibenzothiophenyl group, and a substituted or unsubstituted dibenzosilolyl group, provided that at least one selected from Ar₁ and Ar₂ may each independently be selected from a substituted or unsubstituted fluorenyl group, a substituted or unsubstituted spiro-bifluorenyl group, a substituted or unsubstituted carbazolyl group, a substituted or unsubstituted dibenzofuranyl group, a substi-

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tuted or unsubstituted dibenzothiophenyl group, and a substituted or unsubstituted dibenzosilolyl group; and

at least one substituent of the substituted phenylene group, substituted naphthylene group, substituted fluorenylene group, substituted phenanthrenylene group, substituted anthracenylene group, substituted triphenylenylene group, substituted pyrenylene group, substituted phenyl group, substituted biphenyl group, substituted terphenyl group, substituted naphthyl group, substituted fluorenyl group, substituted phenanthrenyl group, substituted anthracenyl group, substituted triphenylenyl group, substituted pyrenyl group, substituted fluorenyl group, substituted spiro-bifluorenyl group, substituted carbazolyl group, substituted dibenzofuranyl group, substituted dibenzothiophenyl group, and a substituted dibenzosilolyl group may be selected from the group consisting of:

hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, and a C₁-C₂₀ alkoxy group,

a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, and a phosphoric acid group or a salt thereof,

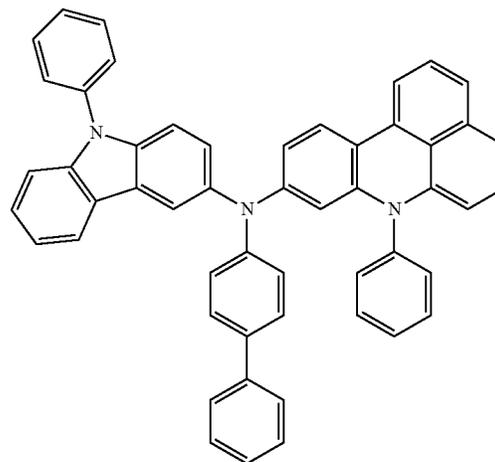
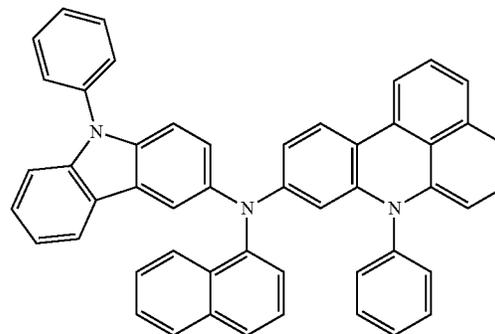
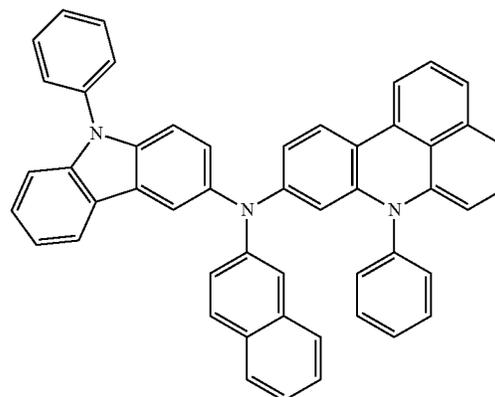
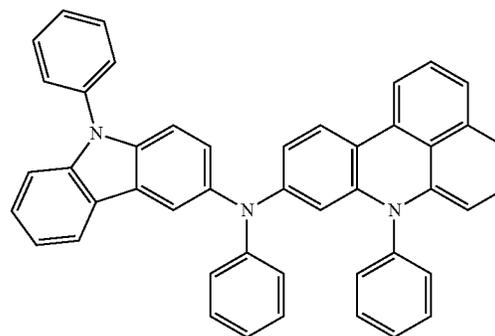
a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a carbazolyl group, a benzocarbazolyl group, and a dibenzocarbazolyl group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a pyridinyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a carbazolyl group, a benzocarbazolyl group, and a dibenzocarbazolyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, and a pyridinyl group, and —Si(Q₃₁)(Q₃₂)(Q₃₃),

wherein Q₃₁ to Q₃₃ may each independently be selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, and a carbazolyl group, but embodiments of the present disclosure are not limited thereto.

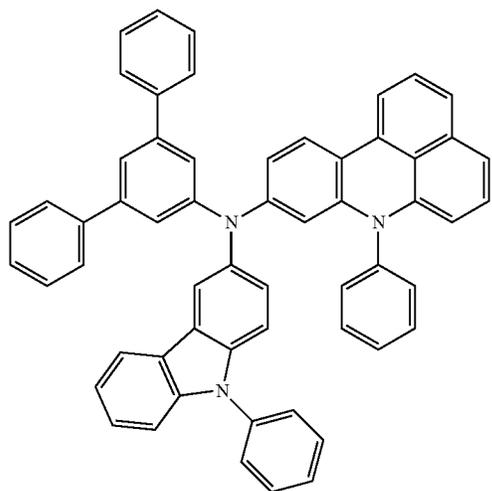
For example, the condensed cyclic compound represented by Formula 1 may be one of Compounds 1 to 96, but is not limited thereto:

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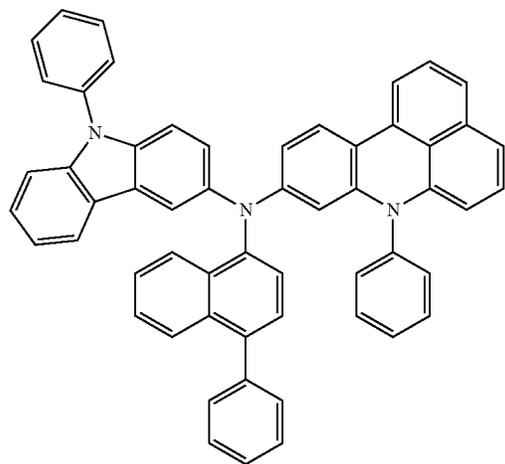
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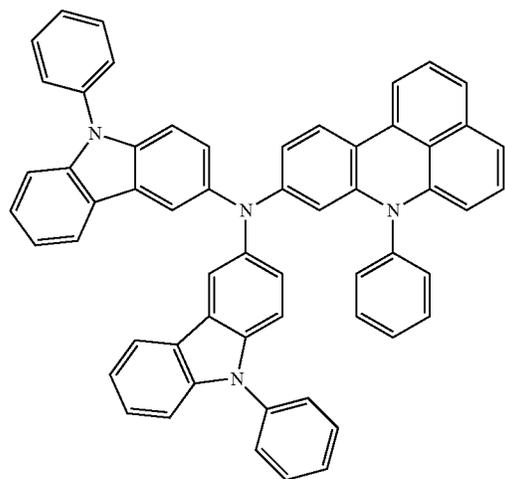
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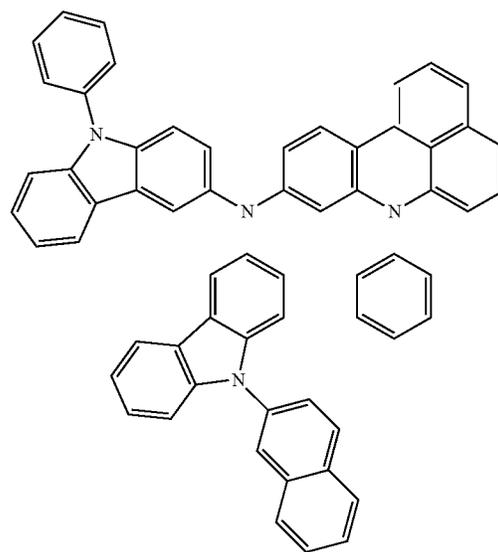
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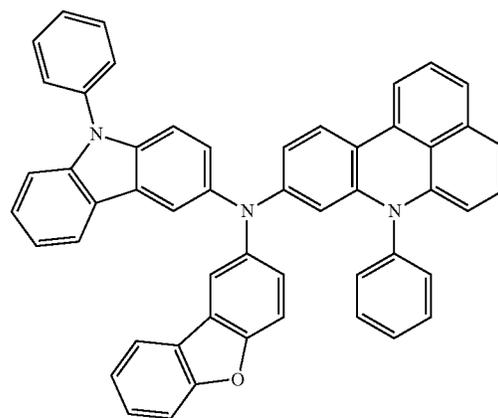
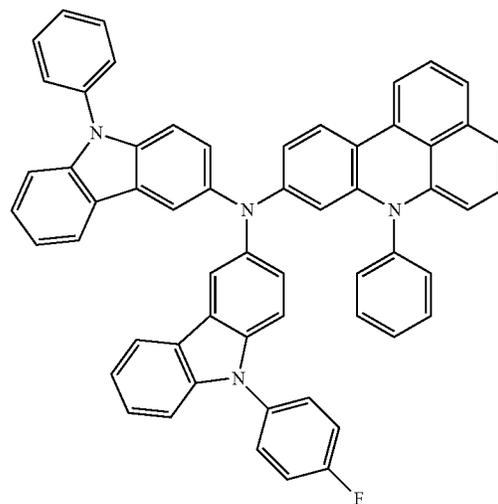
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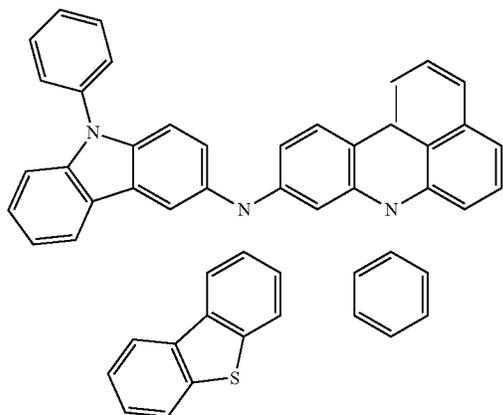
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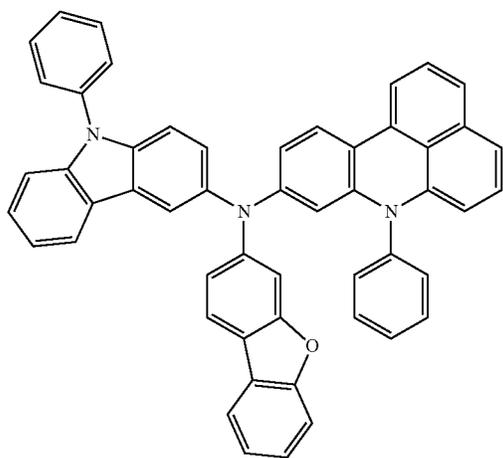
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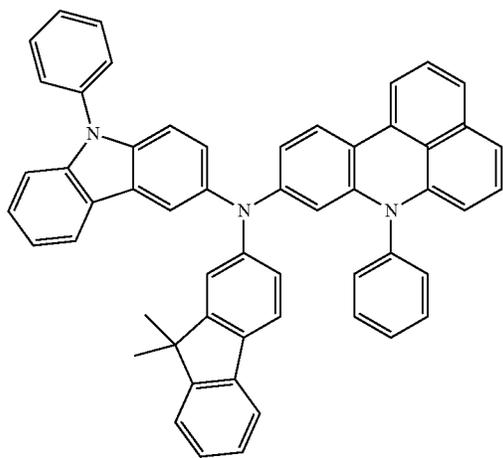
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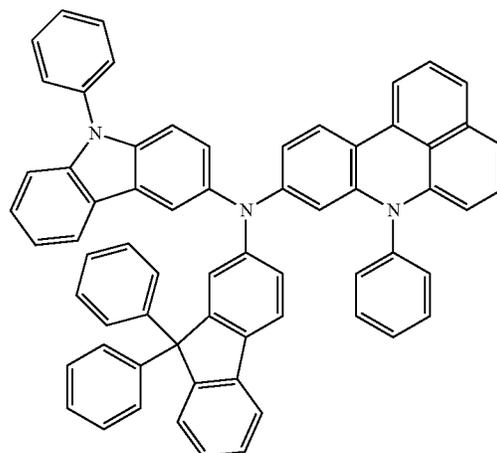
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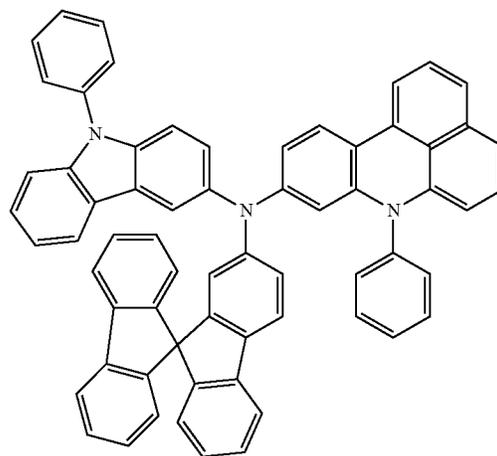
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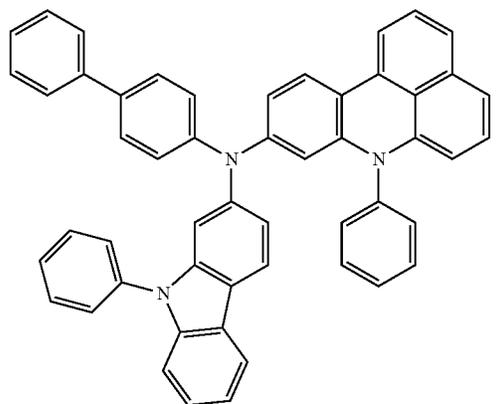
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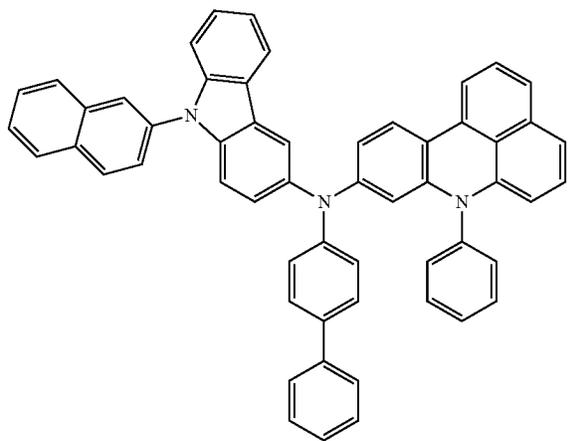
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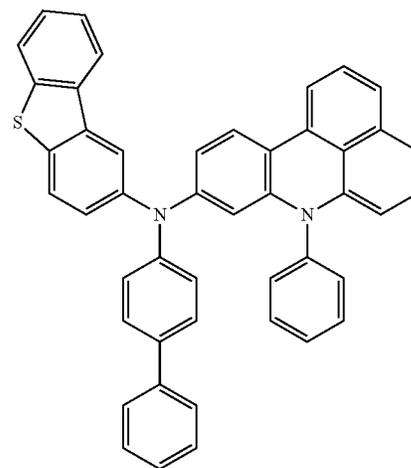
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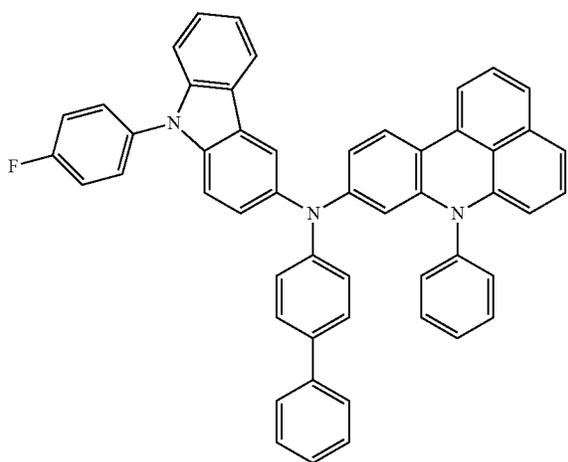
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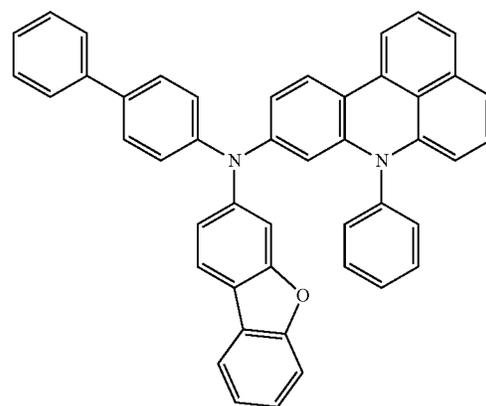


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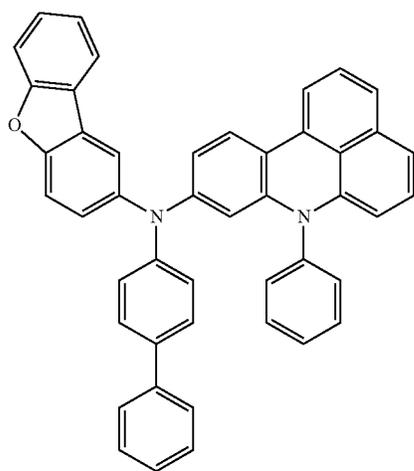
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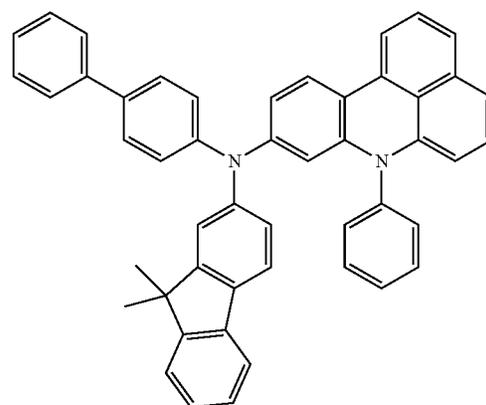
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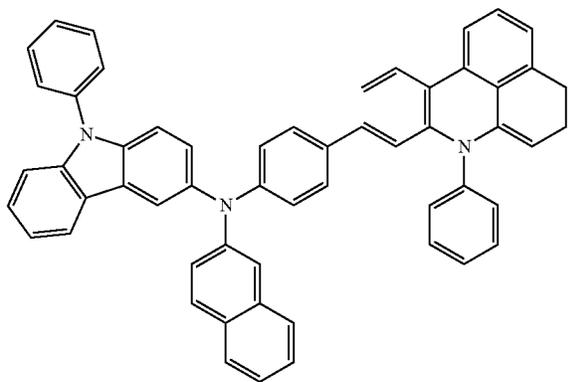
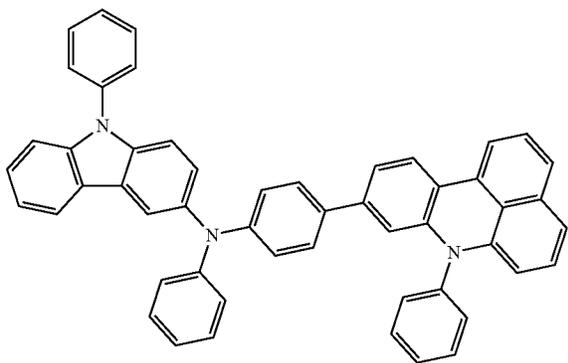
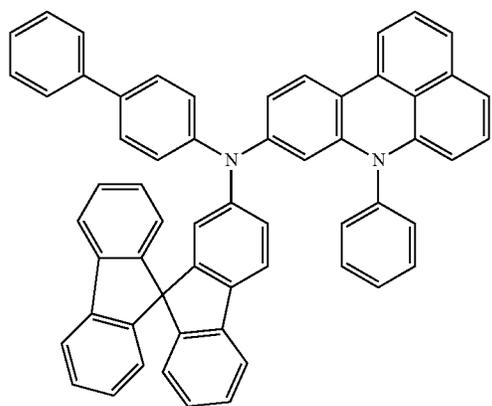
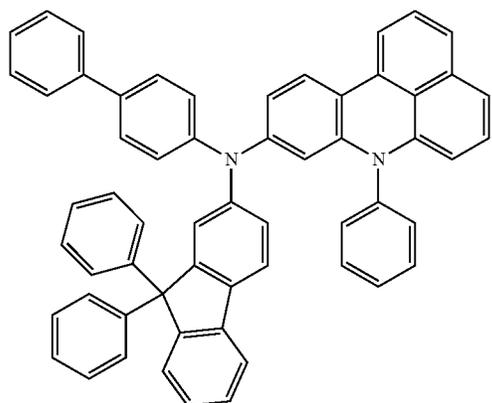
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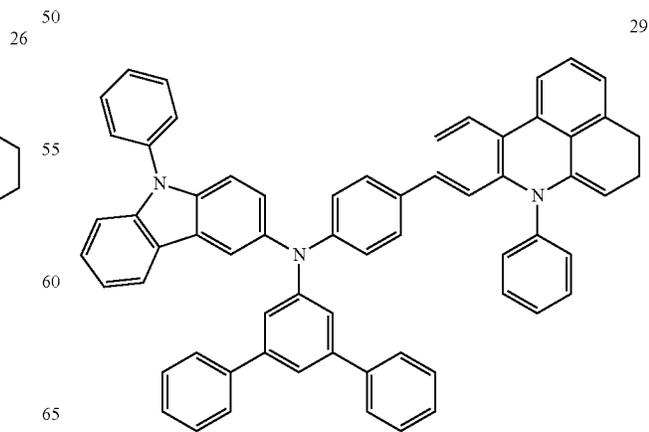
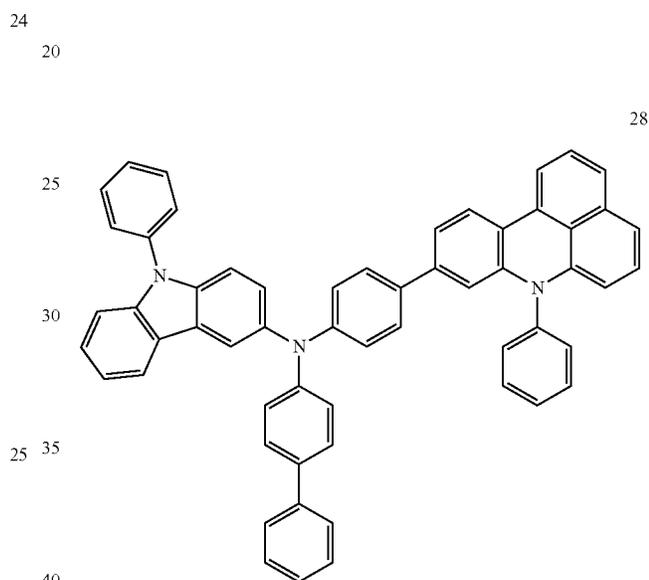
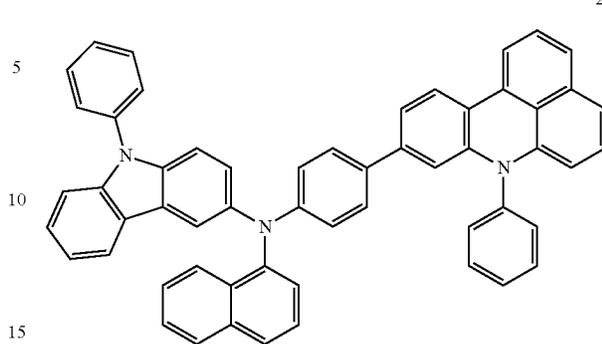


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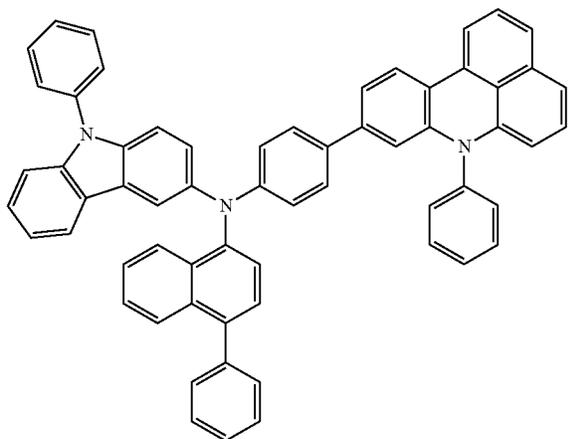
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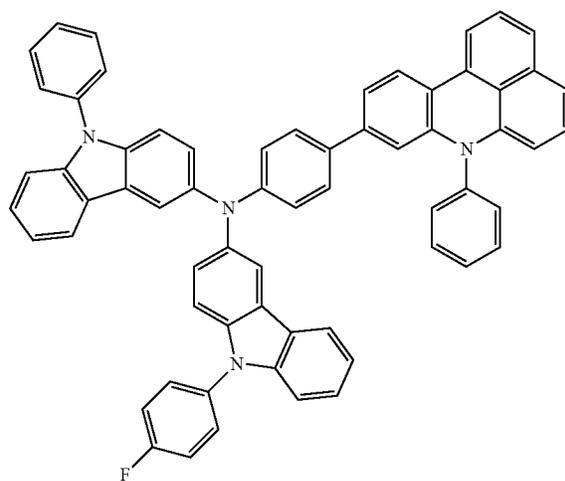
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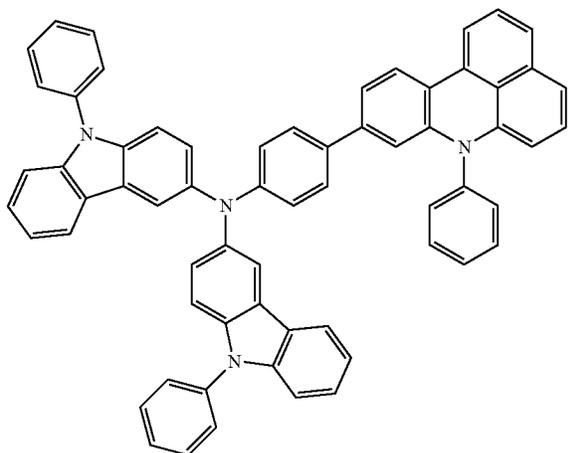
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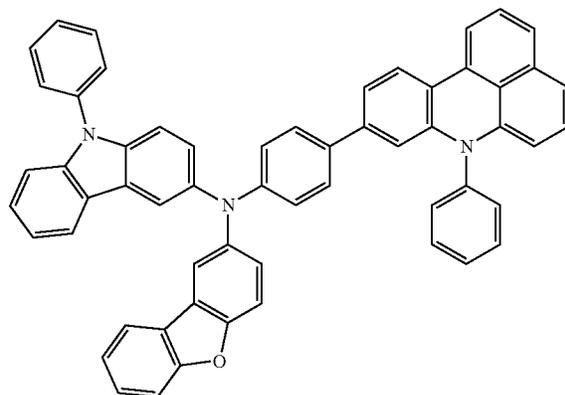


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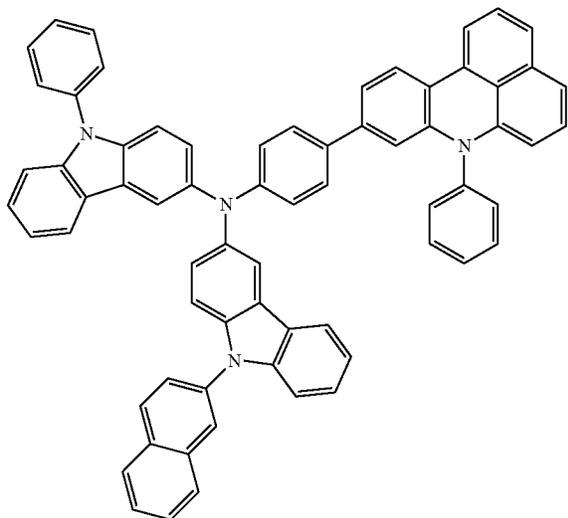
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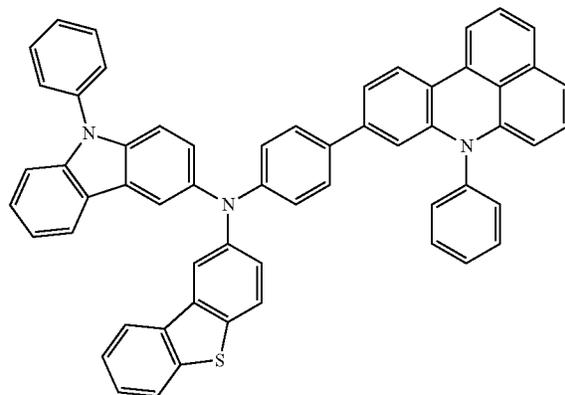
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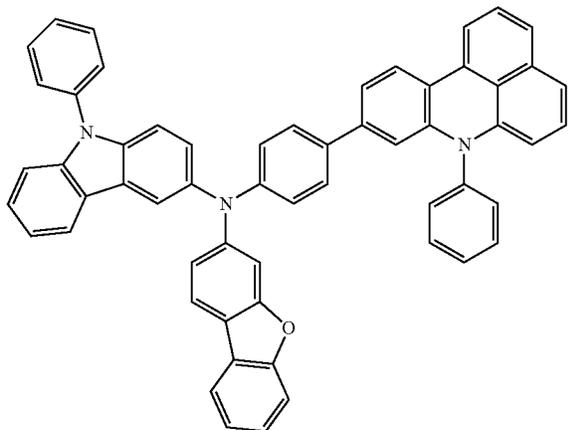
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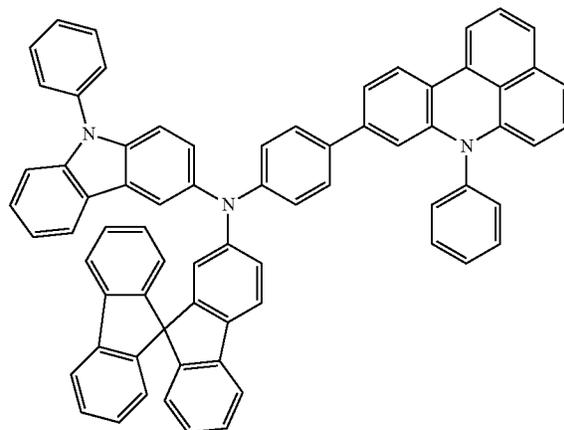
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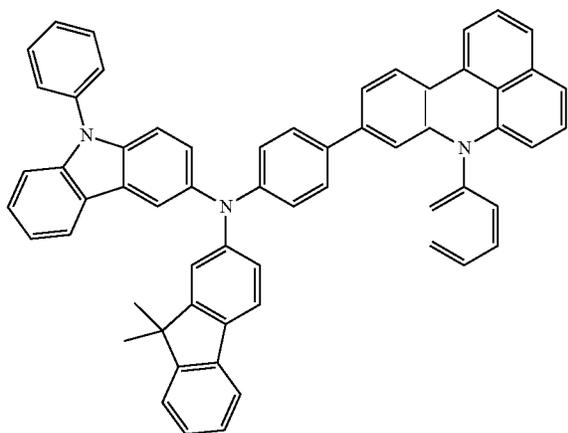
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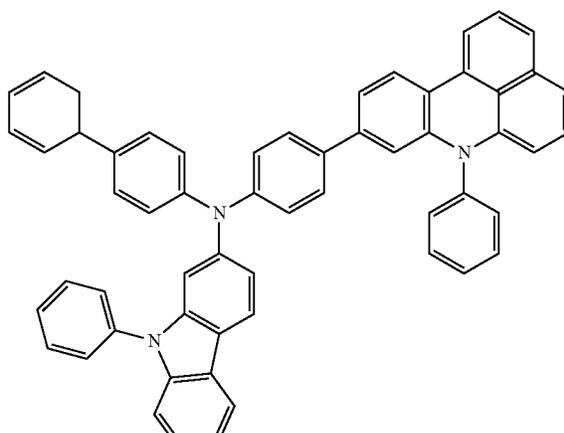
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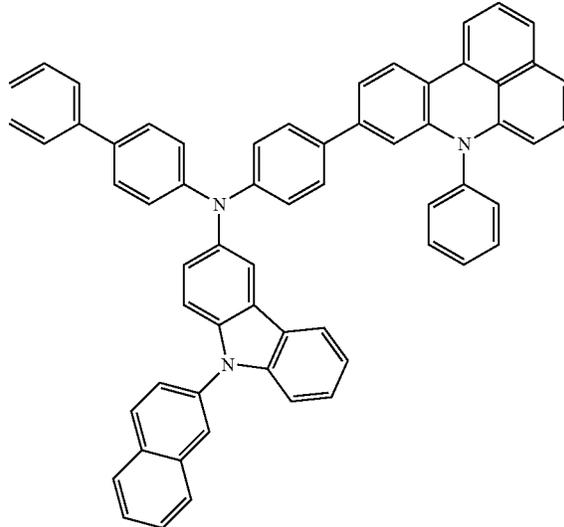
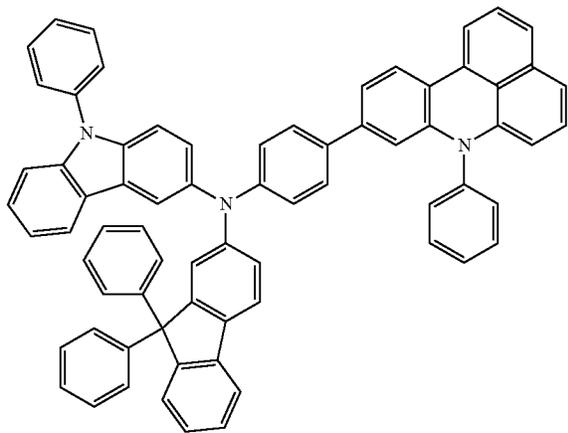
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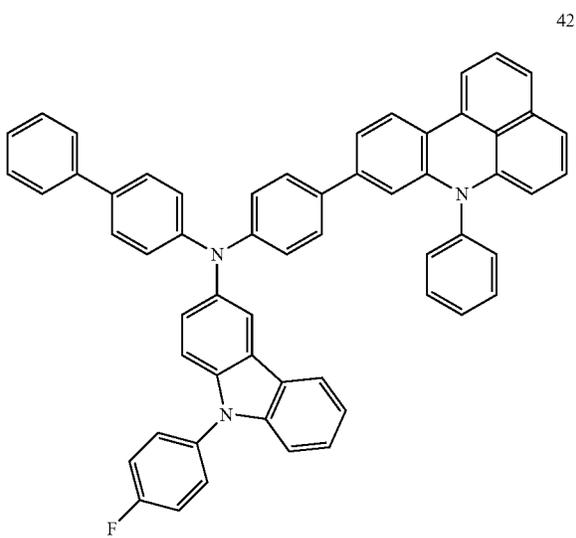
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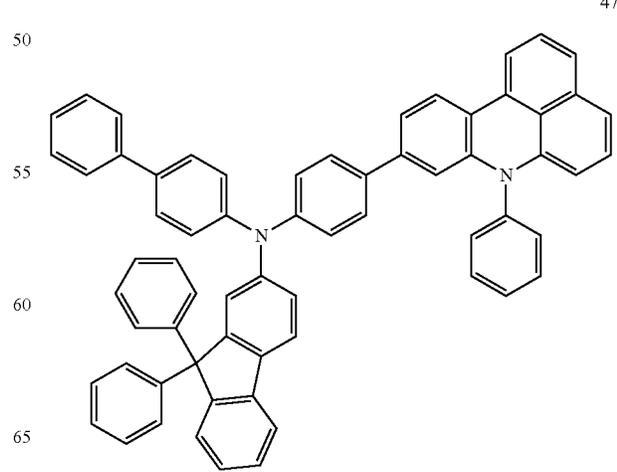
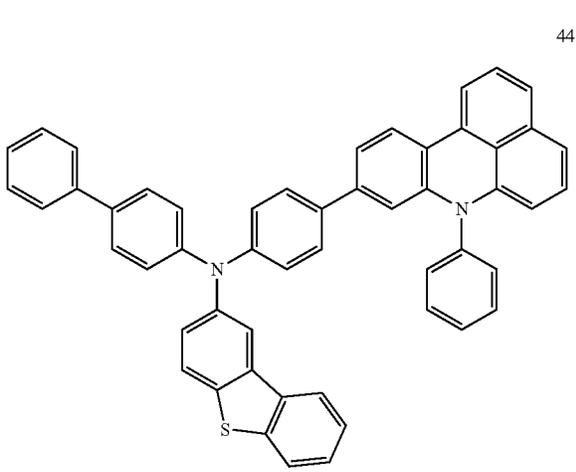
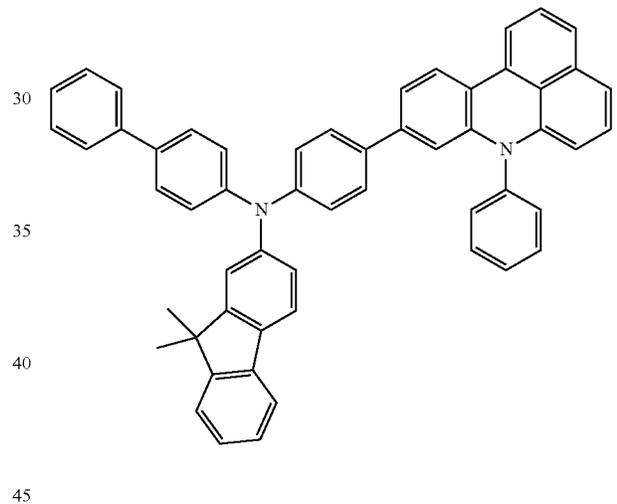
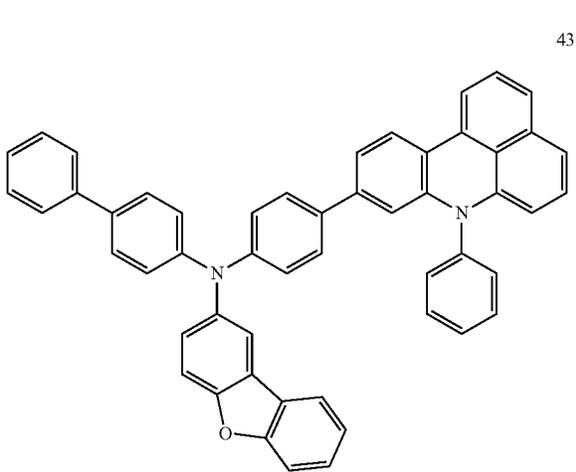
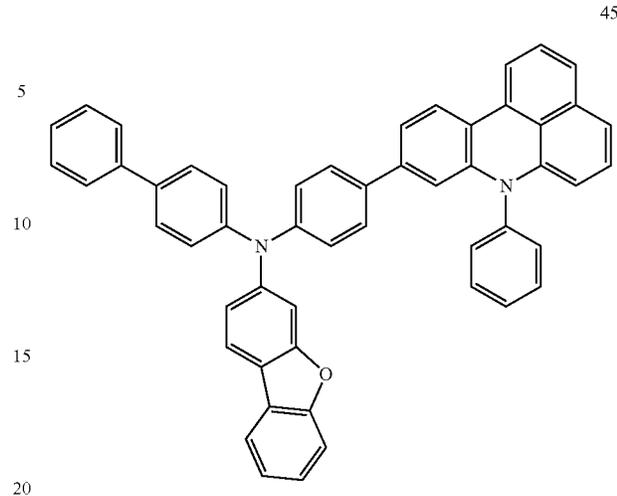
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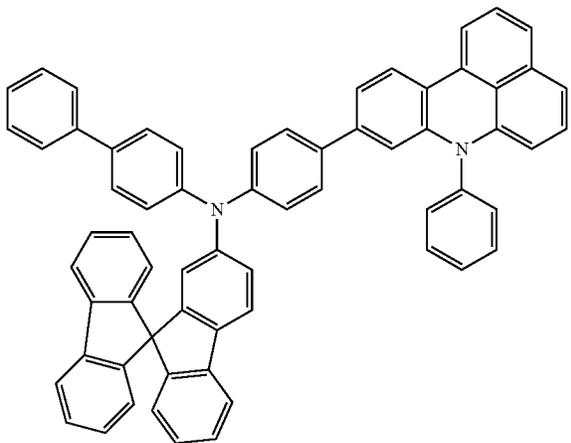
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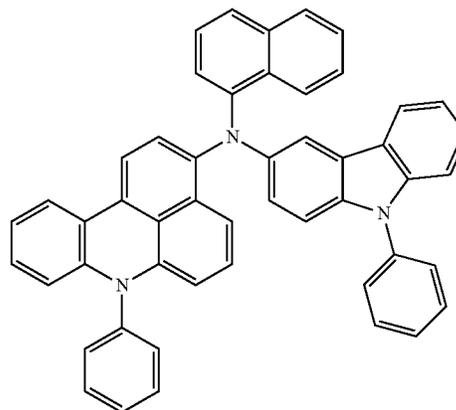
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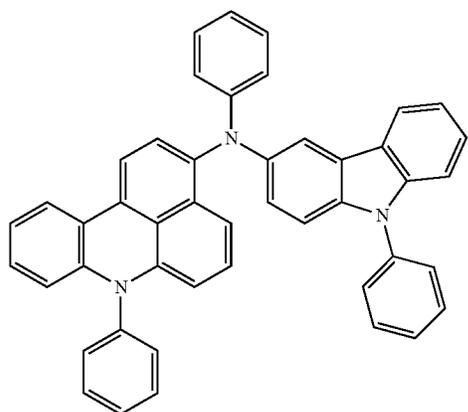
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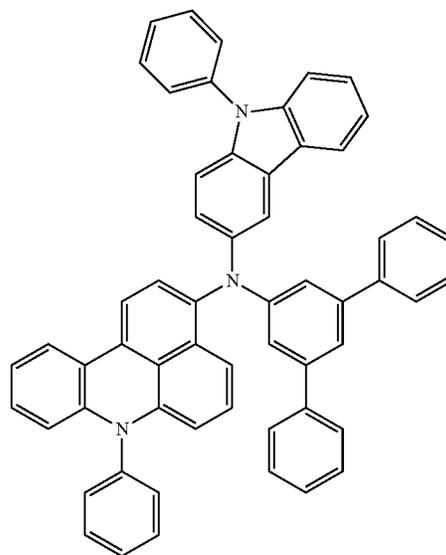
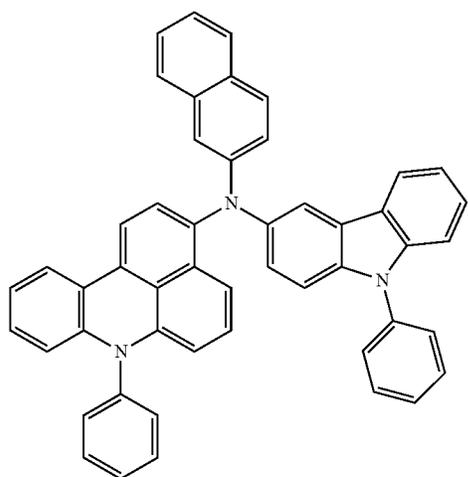
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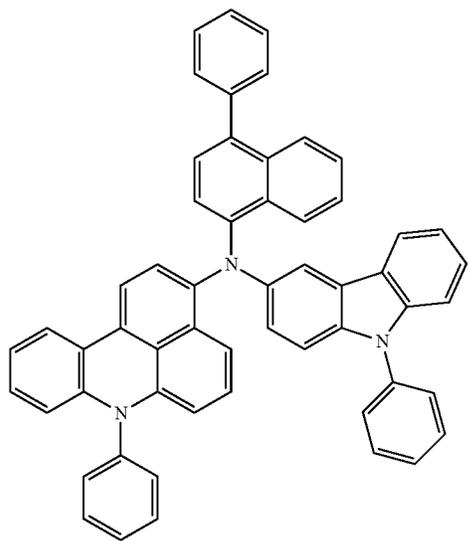
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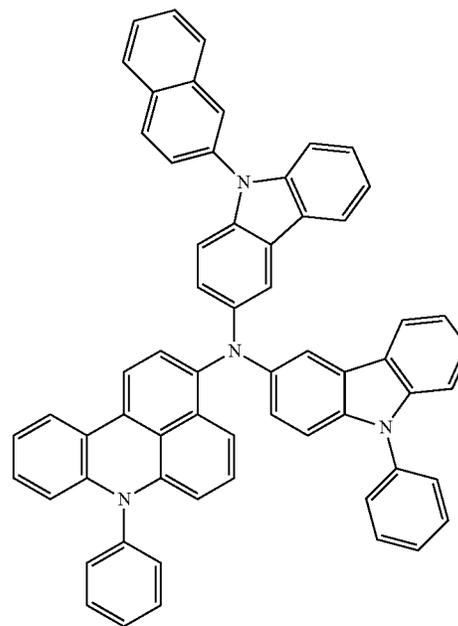
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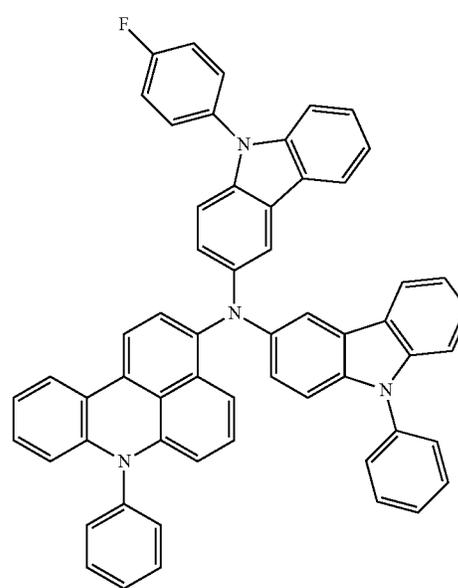
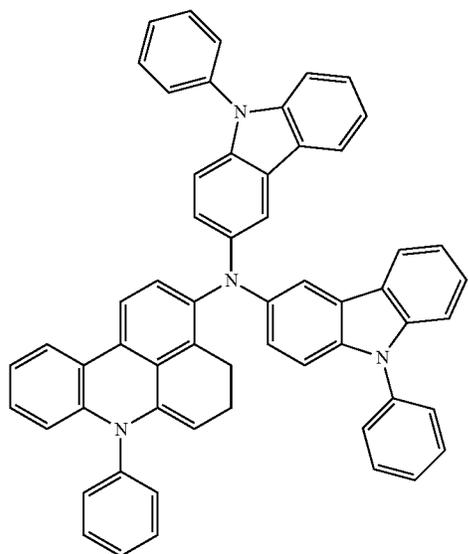
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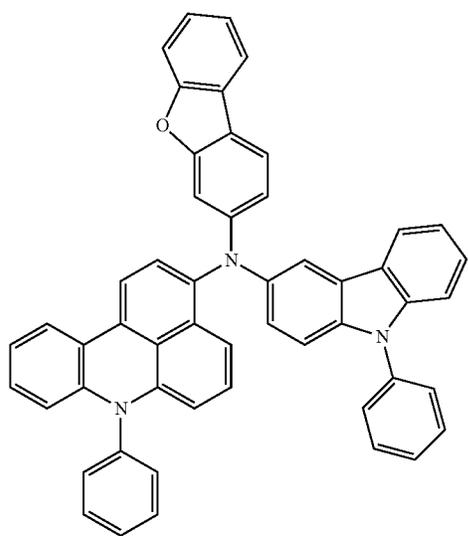
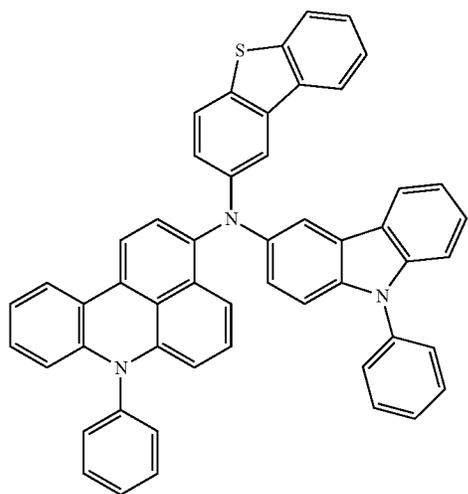
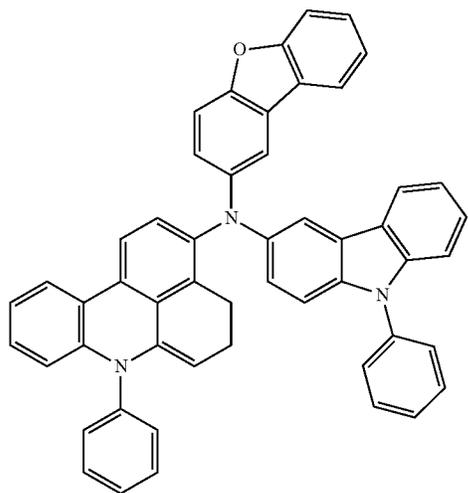
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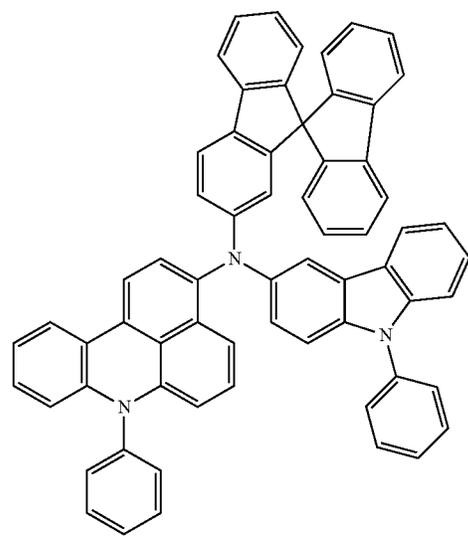
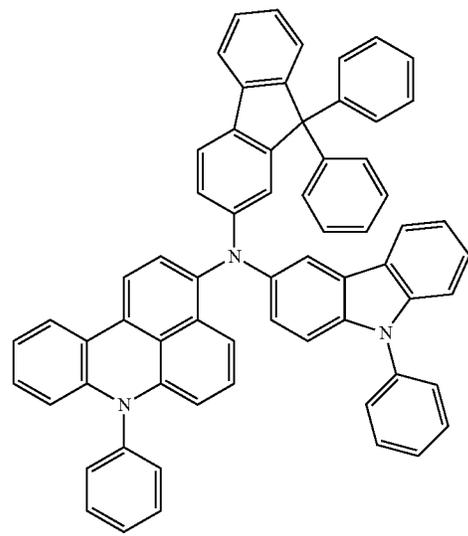
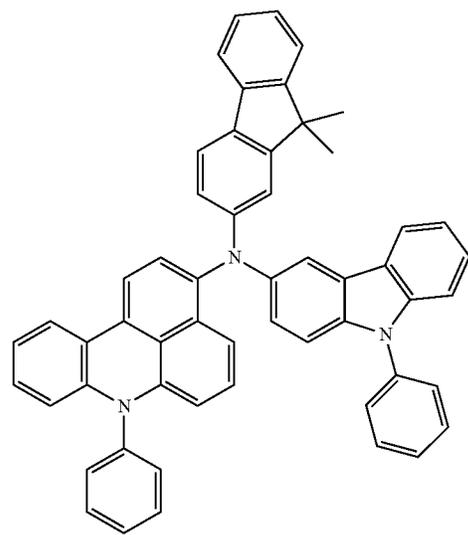
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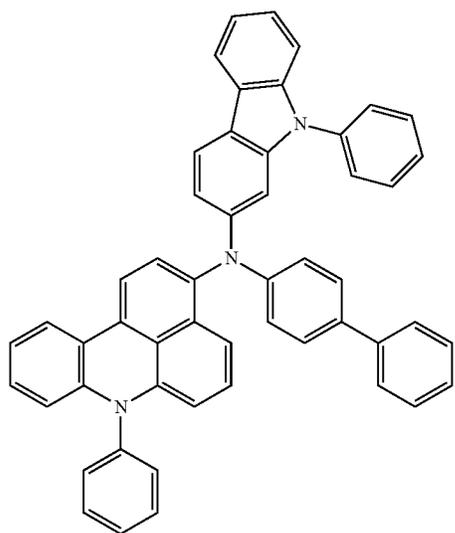
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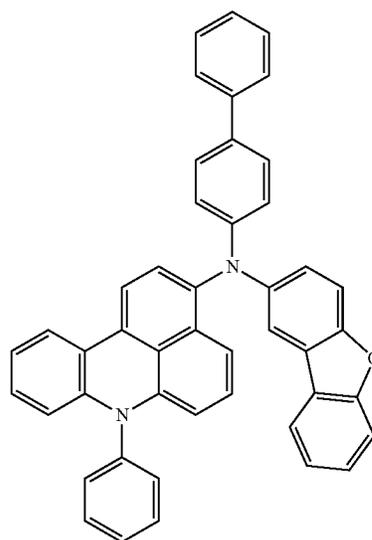
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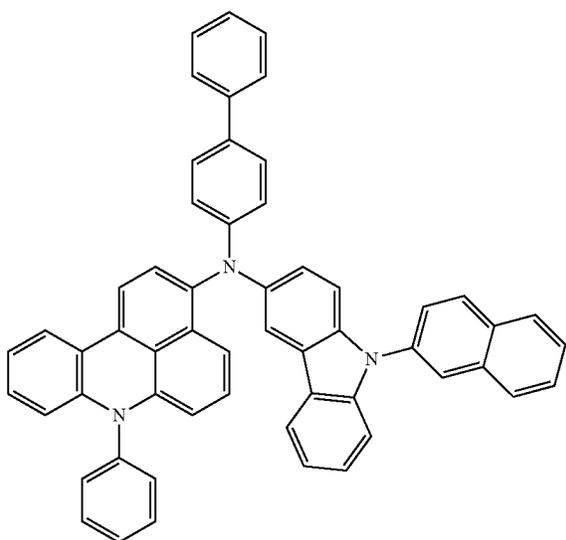
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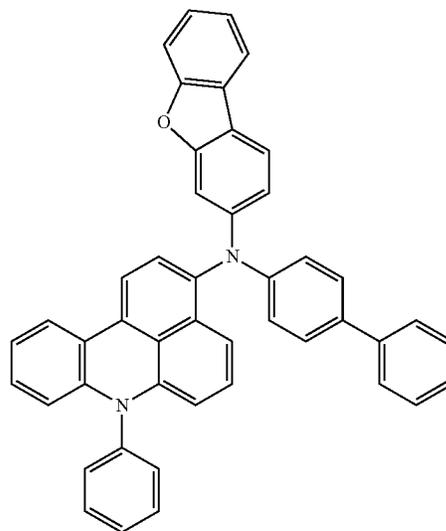
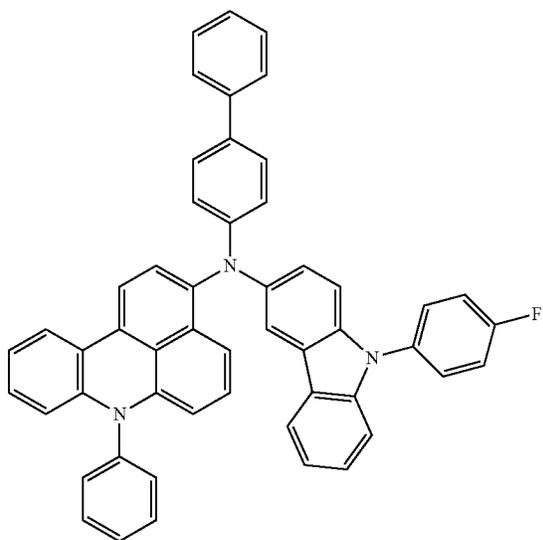
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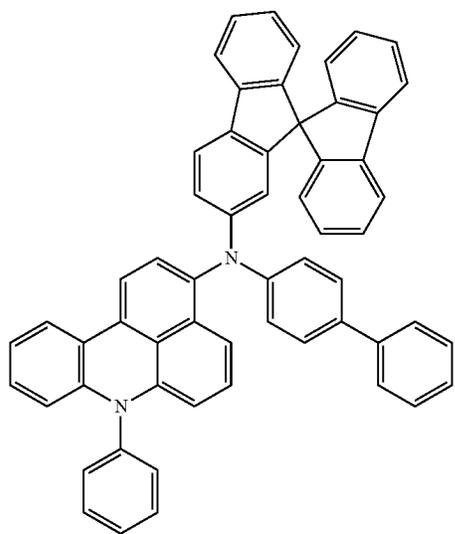
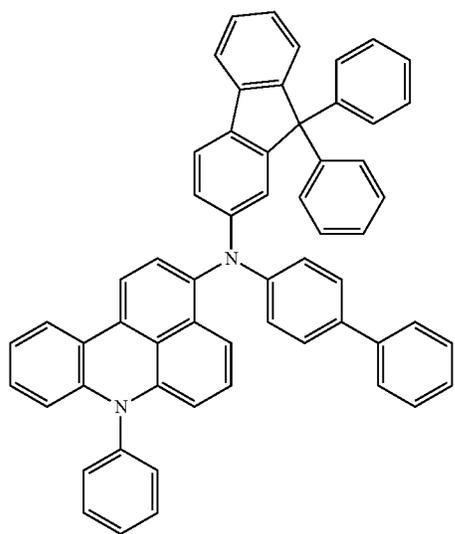
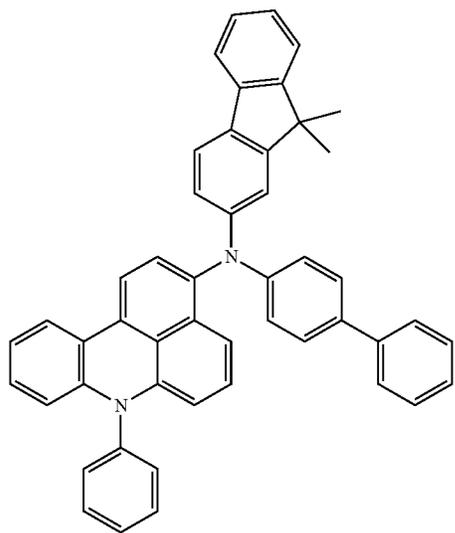
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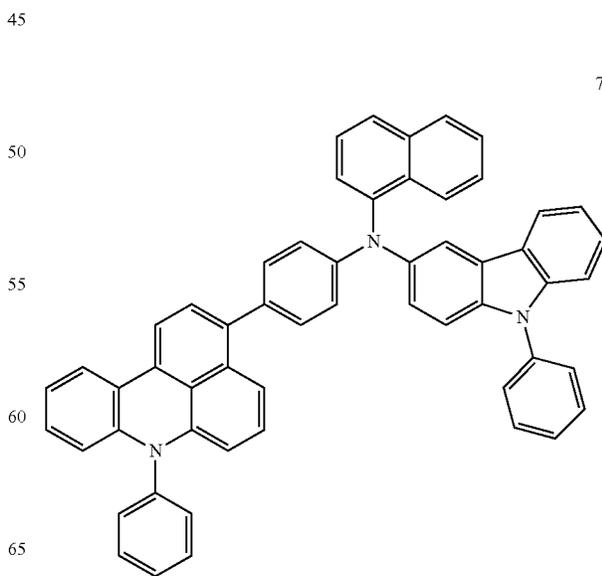
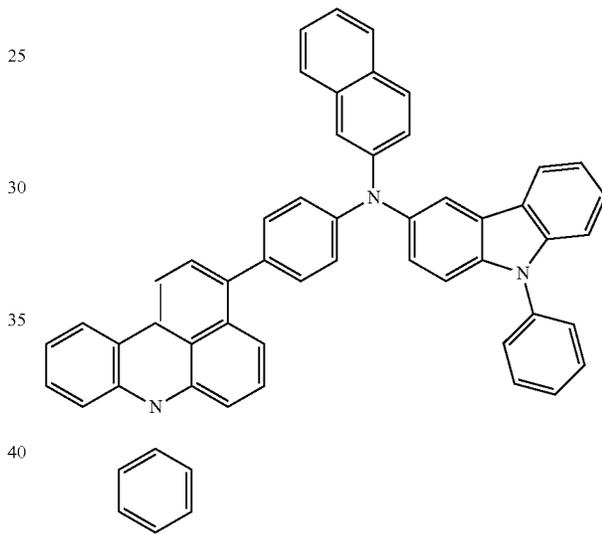
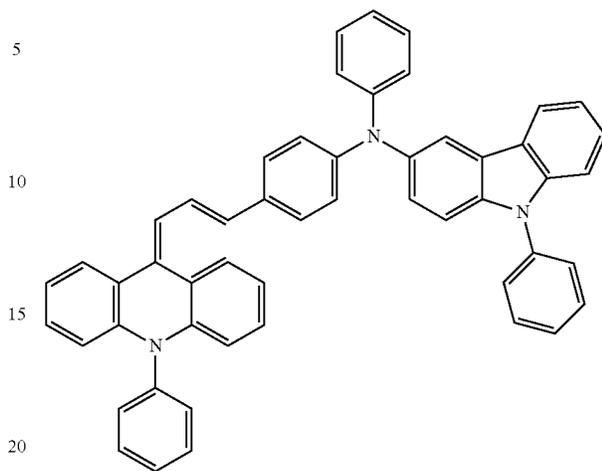
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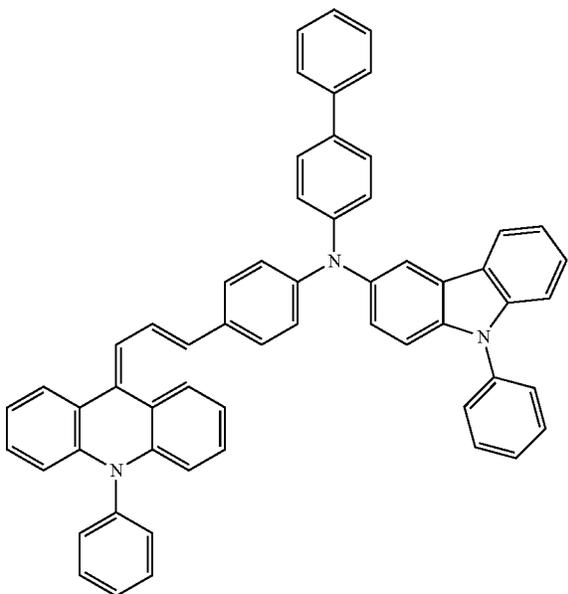
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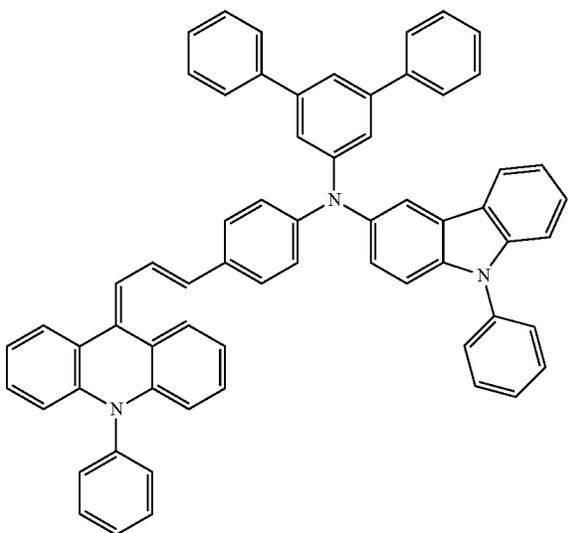
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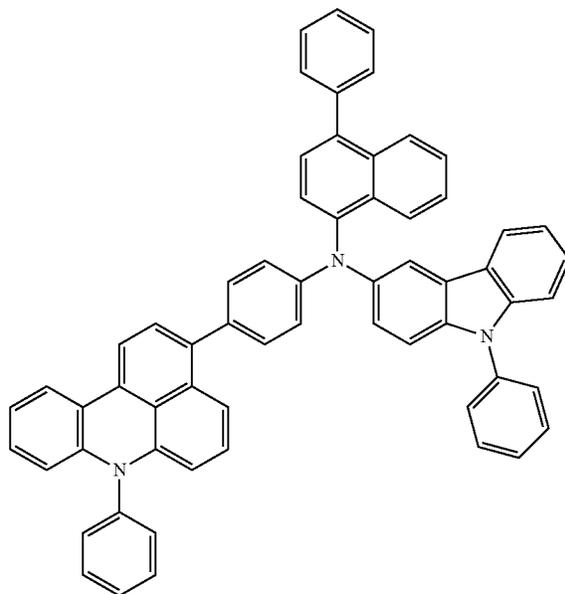
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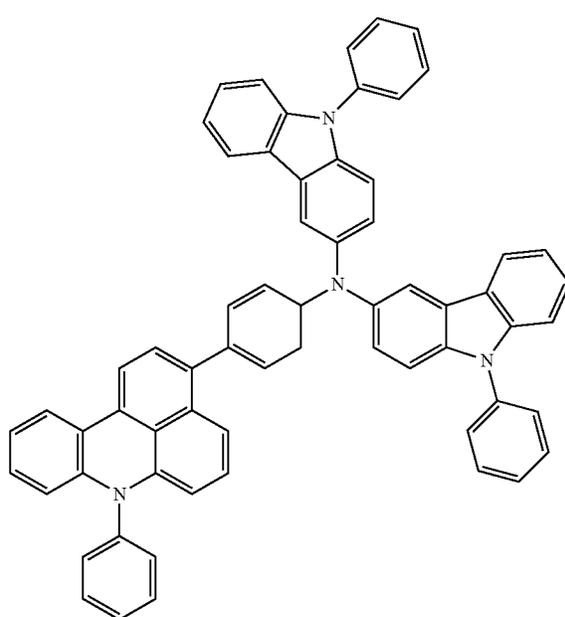
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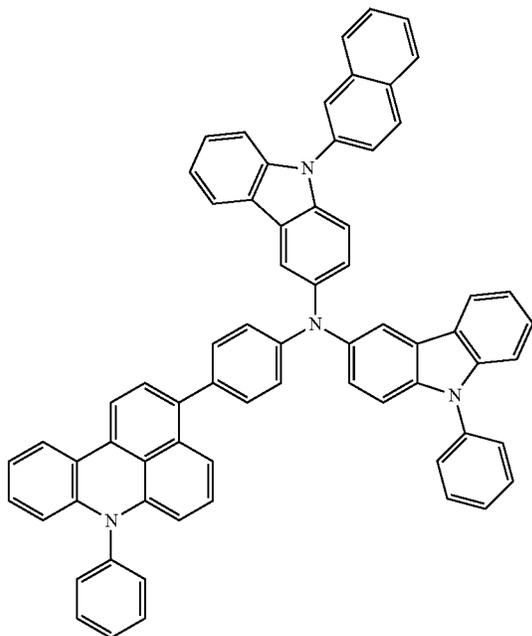
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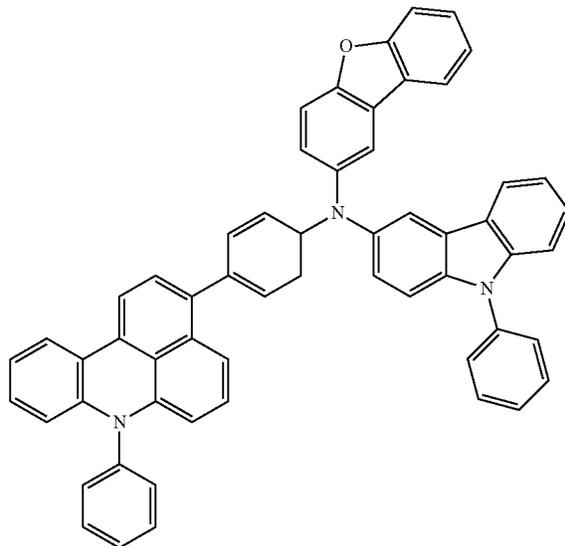
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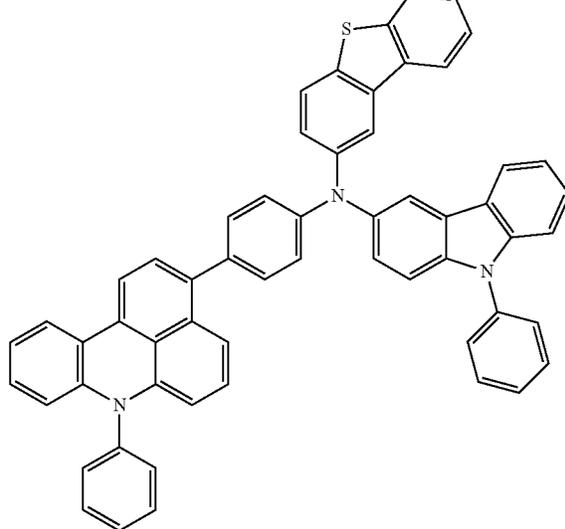
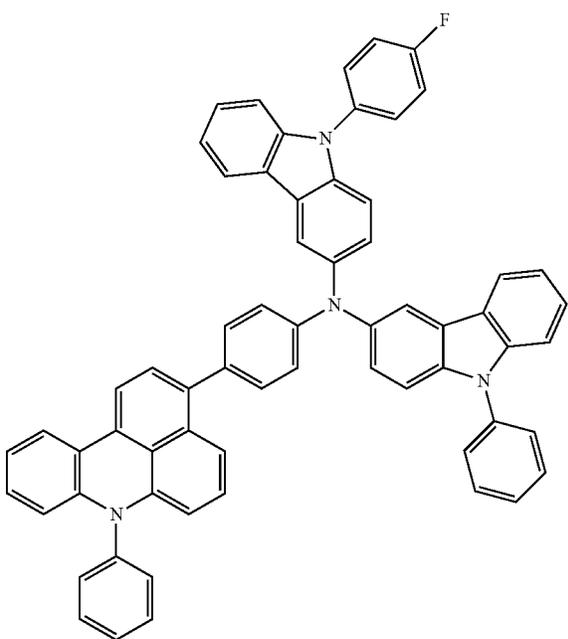
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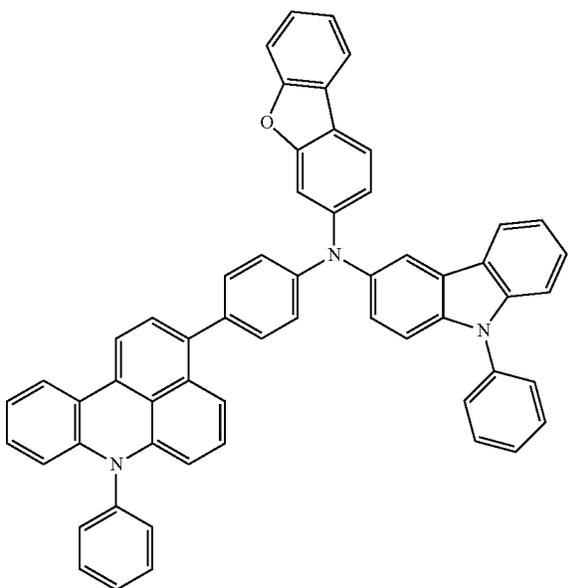
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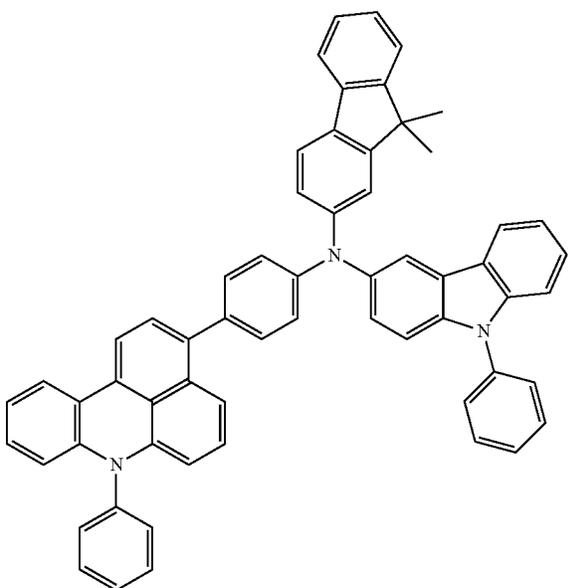
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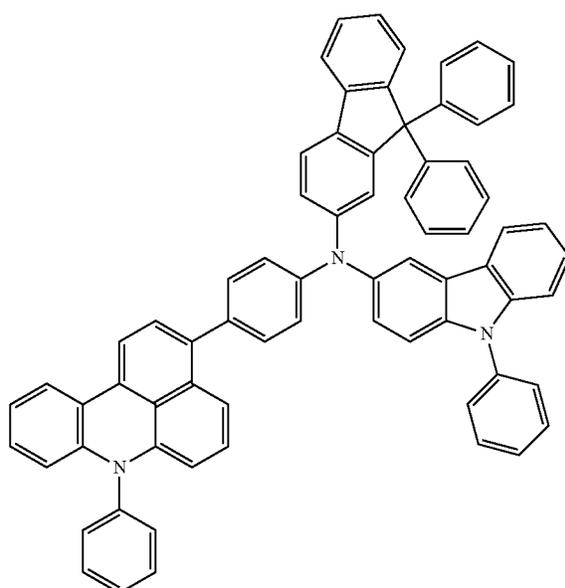
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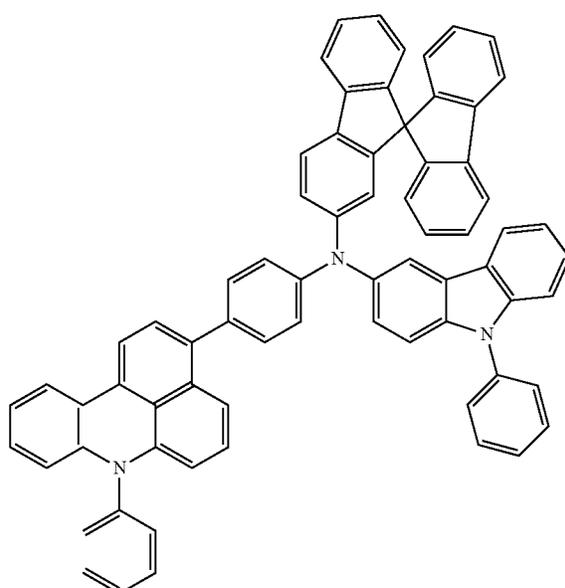
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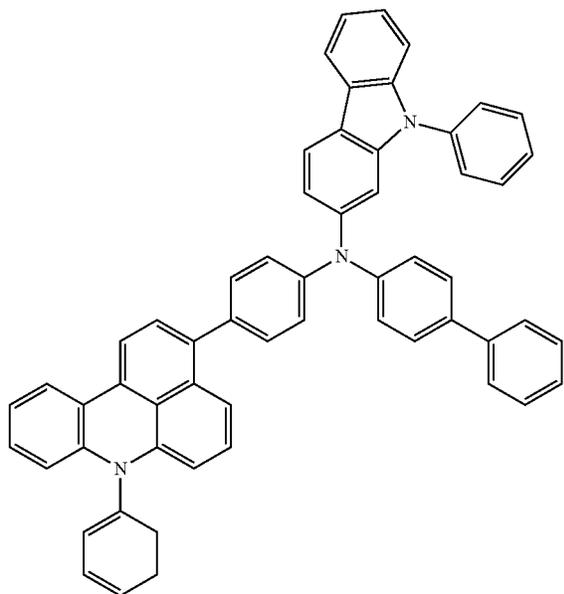
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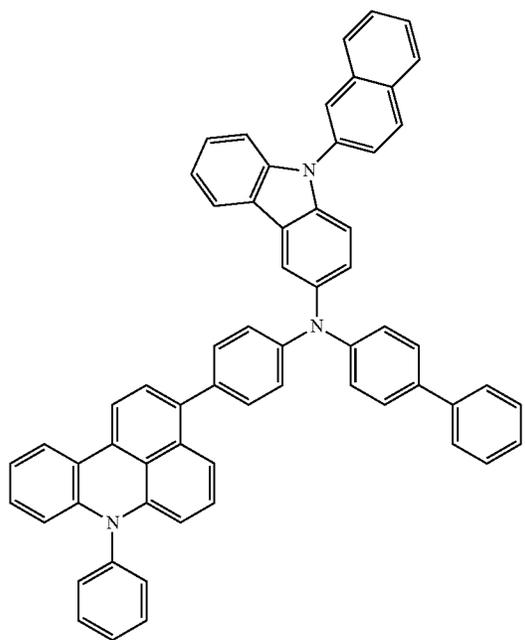
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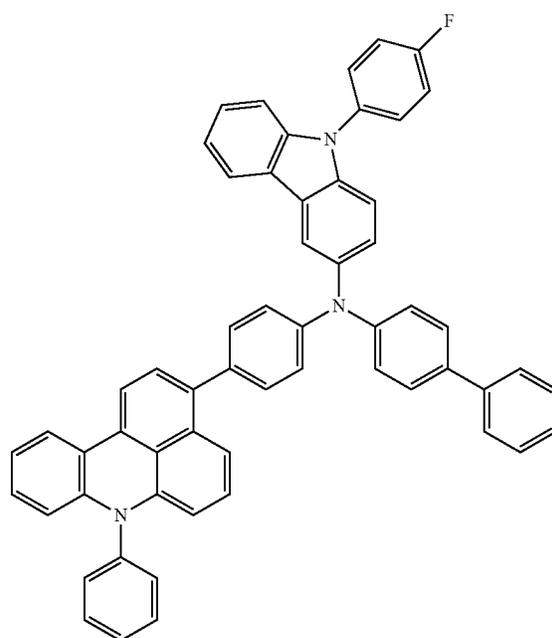
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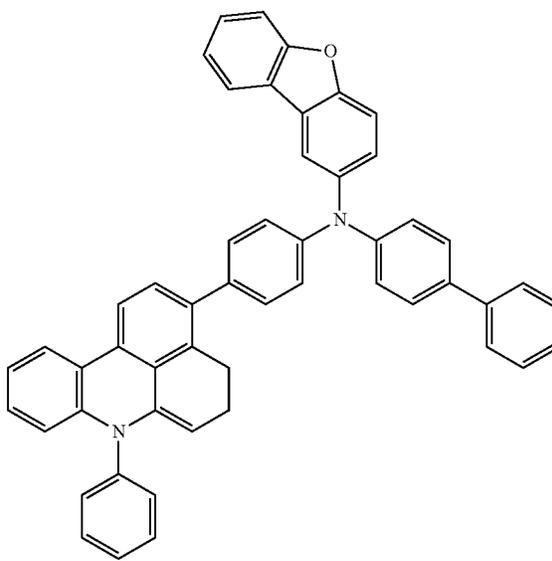
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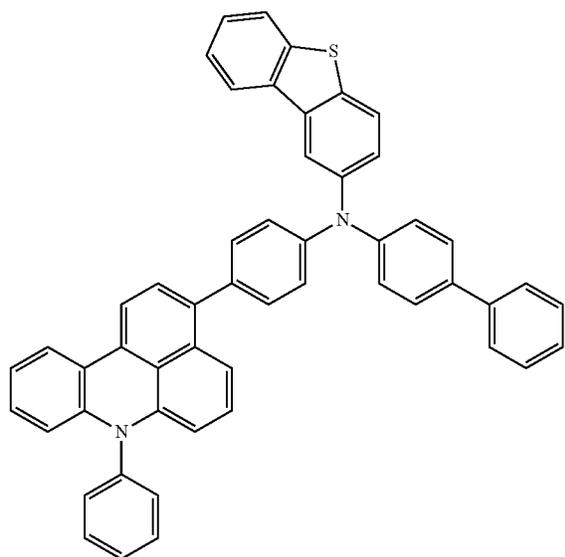


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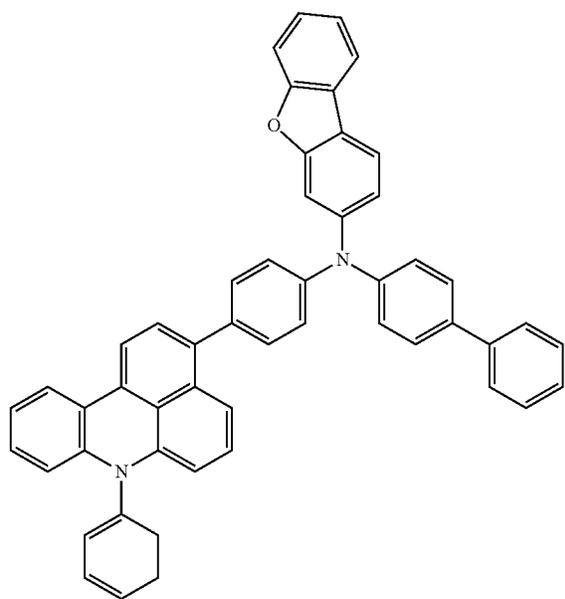
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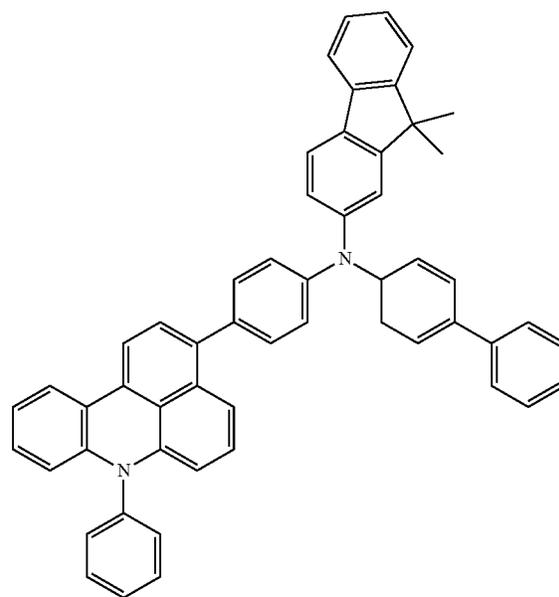
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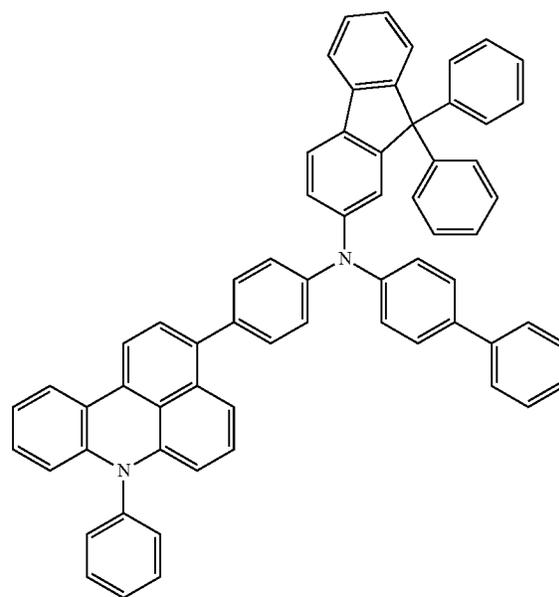


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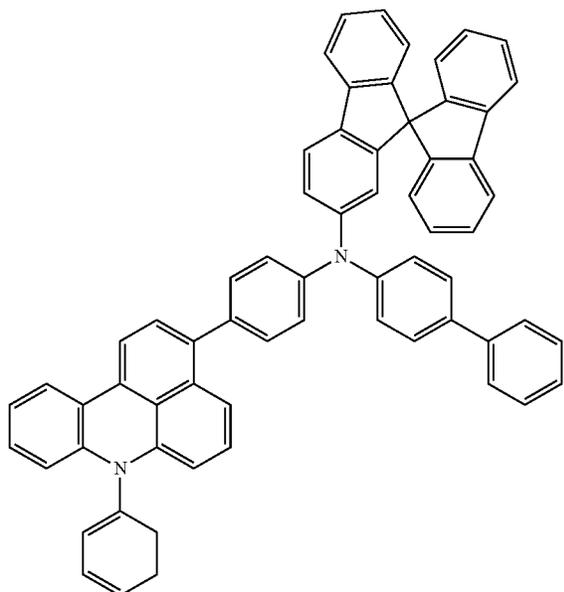
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A substituent selected from R_1 to R_{10} in Formula 1 may be the group represented by Formula 2. That is, the compound represented by Formula 1 may have at least one group represented by Formula 2.

At least one selected from Ar_1 and Ar_2 in Formula 2 may each independently be selected from a substituted or unsubstituted fluorenyl group, a substituted or unsubstituted spiro-bifluorenyl group, a substituted or unsubstituted carbazolyl group, a substituted or unsubstituted dibenzofuranyl group, a substituted or unsubstituted dibenzothiophenyl group, and a substituted or unsubstituted dibenzosilolyl group.

The condensed cyclic compound represented by Formula 1 may have a high glass transition temperature (T_g) or a high melting point. Accordingly, a heat resistance to Joule's heat, which may occur inside an organic layer and/or between the organic layer and an electrode, and durability under high-temperature conditions may increase. Accordingly, an organic light-emitting device including the condensed cyclic compound represented by Formula 1 may show excellent durability during storage and driving. According to embodiments of the present disclosure, the condensed cyclic compound represented by Formula 1 may have a higher melting point than that of a hole transport material of the related art. When deposited, the condensed cyclic compound represented by Formula 1 shows sublimating characteristics, and accordingly, compared to the hole transport material of the related art, may provide processability during a deposition process and an improvement in driving voltage of a device.

The condensed cyclic compound represented by Formula 1 may be synthesized by using one or more suitable organic synthesis methods. Such organic synthesis methods should become apparent to those of ordinary skill in the art by referring to the examples provided below.

At least one condensed cyclic compound represented by Formula 1 may be used between a pair of electrodes of an organic light-emitting device. For example, the condensed cyclic compound may be included in a hole transport region, for example, a hole transport layer. Accordingly, embodiments of the present disclosure provide an organic light-emitting device including: a first electrode; a second elec-

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trode facing the first electrode; and an organic layer between the first electrode and the second electrode, the organic layer including an emission layer, wherein the organic layer includes at least one condensed cyclic compound represented by Formula 1.

The expression "(an organic layer) includes at least one condensed cyclic compound" used herein may include a case in which (an organic layer) includes one or more identical compounds represented by Formula 1 and a case in which (an organic layer) includes two or more different condensed cyclic compounds represented by Formula 1.

For example, the organic layer may include, as the condensed cyclic compound, only Compound 1. In this regard, Compound 1 may be included in an emission layer of the organic light-emitting device. In some embodiments, the organic layer may include, as the condensed cyclic compound, Compound 1 and Compound 2. In this regard, Compound 1 and Compound 2 may both be included in the same layer (e.g., Compound 1 and Compound 2 may both be in an emission layer), or Compound 1 and Compound 2 may be in different layers (e.g., Compound 1 may be in a hole transport layer and Compound 2 may be in an emission layer).

The organic layer may include a hole transport region (between a first electrode (e.g., anode) and an emission layer) and an electron transport region (between the emission layer and a second electrode (e.g., cathode)). The hole transport region may include at least one layer selected from a hole injection layer, a hole transport layer, a buffer layer, and an electron blocking layer; and the electron transport region may include at least one layer selected from a hole blocking layer, an electron transport layer, and an electron injection layer. At least one selected from the hole transport region and the emission layer may include at least one condensed cyclic compound represented by Formula 1. For example, the hole transport region may include the hole transport layer, and the hole transport layer may include at least one condensed cyclic compound represented by Formula 1.

The term "organic layer" used herein may refer to a single layer and/or a plurality of layers disposed (e.g., positioned) between the first electrode and the second electrode of the organic light-emitting device. A material included in the "organic layer" is not limited to an organic material.

The drawing is a schematic view of an organic light-emitting device 10 according to an embodiment. The organic light-emitting device 10 includes a first electrode 110, an organic layer 150, and a second electrode 190.

Hereinafter, the structure of an organic light-emitting device according to an embodiment and a method of manufacturing an organic light-emitting device according to an embodiment will be described in connection with the drawing.

In the drawing, a substrate may be additionally disposed under the first electrode 110 or above the second electrode 190. The substrate may be a glass substrate and/or a transparent plastic substrate, each having excellent mechanical strength, thermal stability, transparency, surface smoothness, ease of handling, and/or water-resistance.

The first electrode 110 may be formed by depositing or sputtering a material for forming the first electrode 110 on the substrate. When the first electrode 110 is an anode, for example, the material for forming the first electrode 110 may be selected from materials with a high work function, so as to facilitate hole injection. The first electrode 110 may be a reflective electrode, a semi-transmissive electrode, or a transmissive electrode. The material for forming the first

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electrode **110** may be a transparent and highly conductive material, and non-limiting examples of such material include indium tin oxide (ITO), indium zinc oxide (IZO), tin oxide (SnO₂), and zinc oxide (ZnO). When the first electrode **110** is a semi-transmissive electrode or a reflective electrode, at least one selected from magnesium (Mg), aluminum (Al), aluminum-lithium (Al—Li), calcium (Ca), magnesium-indium (Mg—In), and magnesium-silver (Mg—Ag) may be used as a material for forming the first electrode **110**.

The first electrode **110** may have a single-layer structure, or a multi-layer structure including two or more layers. For example, the first electrode **110** may have a three-layered structure of ITO/Ag/ITO, but the structure of the first electrode **110** is not limited thereto.

The organic layer **150** may be disposed on the first electrode **110**. The organic layer **150** may include an emission layer.

The organic layer **150** may further include a hole transport region between the first electrode and the emission layer, and an electron transport region between the emission layer and the second electrode.

The hole transport region may include at least one selected from a hole injection layer (HIL), a hole transport layer (HTL), a buffer layer, and an electron blocking layer (EBL); and the electron transport region may include at least one selected from a hole blocking layer (HBL), an electron transport layer (ETL), and an electron injection layer (EIL), but they are not limited thereto.

The hole transport region may have a single-layered structure formed of a single material, a single-layered structure formed of a plurality of different materials, or a multi-layered structure having a plurality of layers formed of a plurality of different materials.

For example, the hole transport region may have a single-layered structure formed of a plurality of different materials, or a structure of hole injection layer/hole transport layer, a structure of hole injection layer/hole transport layer/buffer layer, a structure of hole injection layer/buffer layer, a structure of hole transport layer/buffer layer, or a structure of hole injection layer/hole transport layer/electron blocking layer, wherein the layers of each structure are sequentially stacked from the first electrode **110** in this stated order, but the structure of the hole transport region is not limited thereto.

When the hole transport region includes a hole injection layer, the hole injection layer may be formed on the first electrode **110** by using one or more suitable methods such as vacuum deposition, spin coating, casting, a Langmuir-Blodgett (LB) method, ink-jet printing, laser-printing, and/or laser-induced thermal imaging.

When a hole injection layer is formed by vacuum deposition, for example, the vacuum deposition may be performed at a deposition temperature of about 100 to about 500° C., at a vacuum degree of about 10⁻⁸ to about 10⁻³ torr, and at a deposition rate of about 0.01 to about 100 Å/sec, by taking into account a compound for forming the hole injection layer to be deposited, and the structure of the hole injection layer to be formed.

When a hole injection layer is formed by spin coating, for example, the spin coating may be performed at a coating rate of about 2000 rpm to about 5000 rpm, and at a temperature of about 80° C. to 200° C., by taking into account a compound for forming the hole injection layer to be deposited, and the structure of the hole injection layer to be formed.

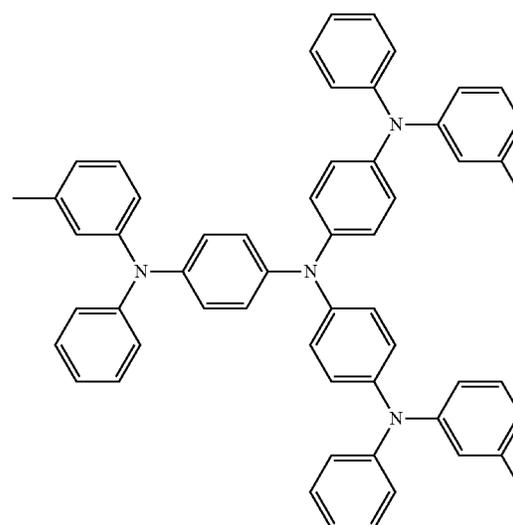
When the hole transport region includes a hole transport layer, the hole transport layer may be formed on the first electrode **110** or the hole injection layer by using one or more suitable methods such as vacuum deposition, spin coating, casting, a LB method, ink-jet printing, laser-print-

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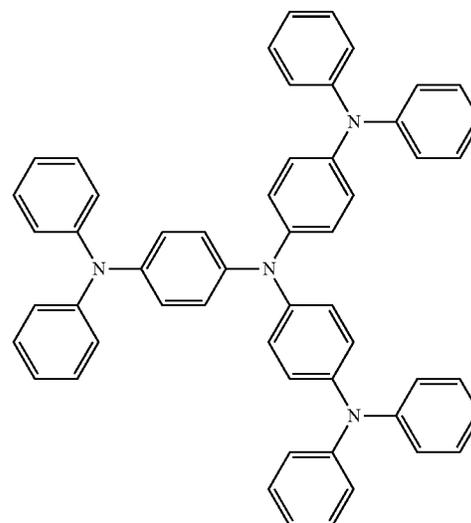
ing, and/or laser-induced thermal imaging. When the hole transport layer is formed by vacuum deposition and/or spin coating, deposition and coating conditions for the hole transport layer may be the same as (or similar to) the deposition and coating conditions for the hole injection layer.

The hole transport region may include the condensed cyclic compound represented by Formula 1. For example, the hole transport region may include the hole transport layer, and the hole transport layer may include the condensed cyclic compound represented by Formula 1.

In various embodiments, the hole transport region may include at least one selected from m-MTDATA, TDATA, 2-TNATA, NPB, β-NPB, TPD, Spiro-TPD, Spiro-NPB, methylated NPB, TAPC, HMTPD, 4,4',4"-tris(N-carbazolyl) triphenylamine (TCTA), polyaniline/dodecylbenzenesulfonic acid (Pani/DBSA), poly(3,4-ethylenedioxythiophene)/poly(4-styrenesulfonate) (PEDOT/PSS), polyaniline/camphor sulfonic acid (Pani/CSA), (polyaniline)/poly(4-styrenesulfonate) (PANI/PSS), a compound represented by Formula 201, and a compound represented by Formula 202:



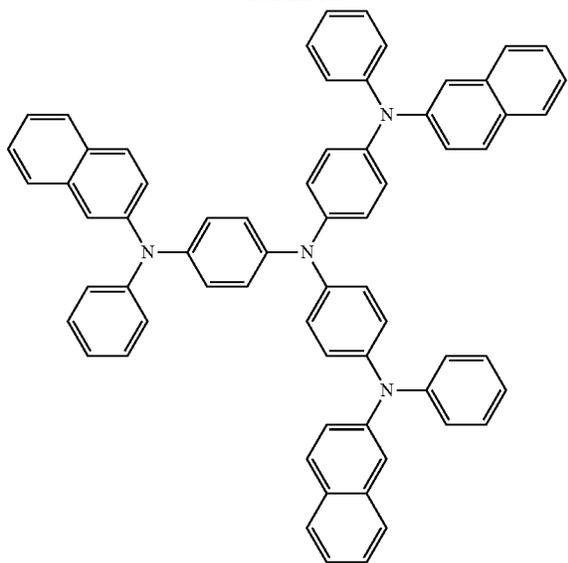
m-MTDATA



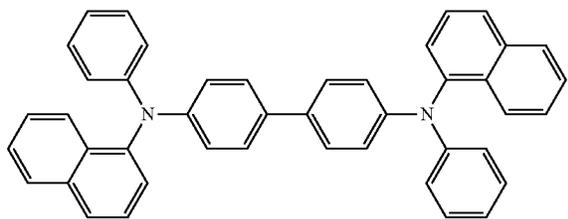
TDATA

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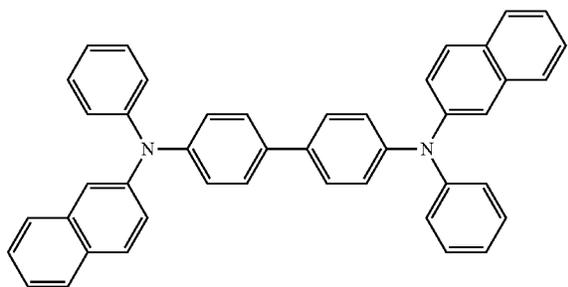
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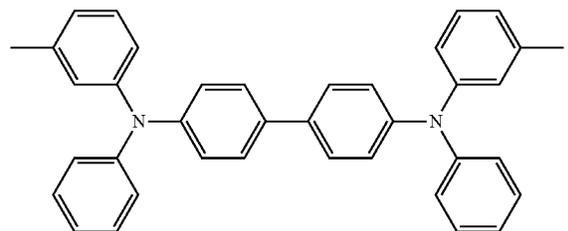
2-TNATA



NPB



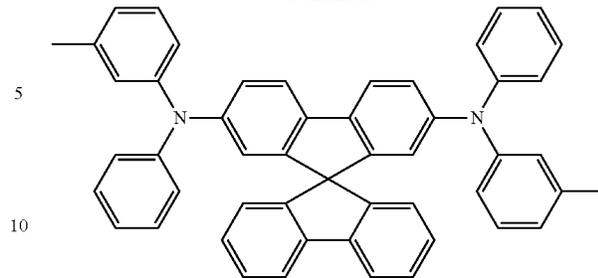
β -NPB



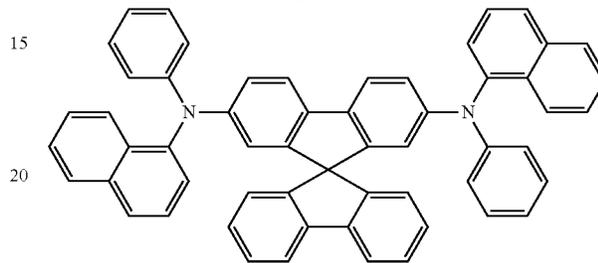
TPD

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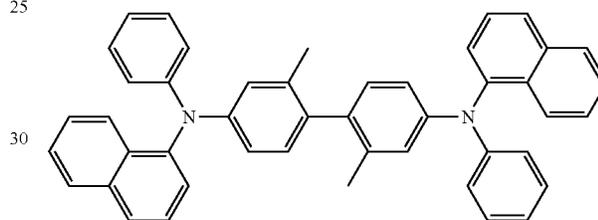
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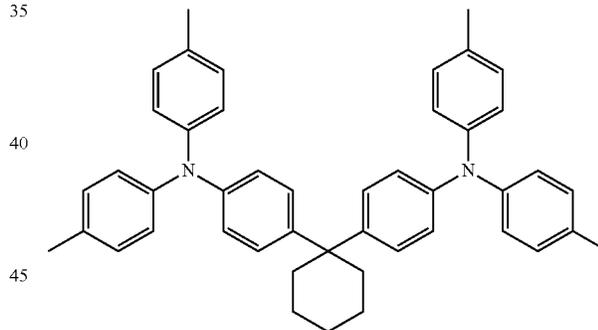
Spiro-TPD



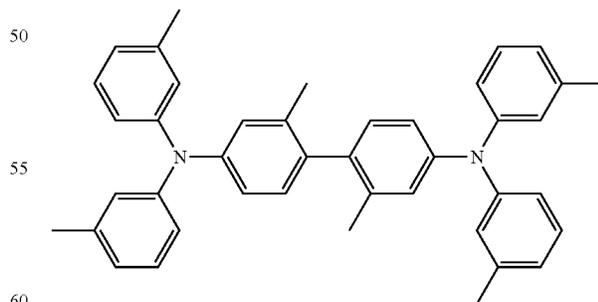
Spiro-NPB



methylated NPB

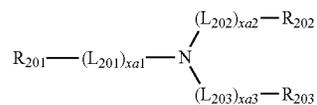


TAPC



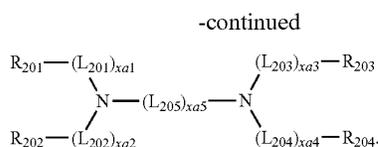
HMTPD

Formula 201



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Formula 202

In Formulae 201 and 202,

Descriptions of L_{201} to L_{205} may each independently be the same as the description provided in connection with L_1 ; $xa1$ to $xa4$ may each independently be selected from 0, 1, 2, and 3;

$xa5$ may be selected from 1, 2, 3, 4, and 5; and

R_{201} to R_{204} may each independently be selected from a substituted or unsubstituted C_3 - C_{10} cycloalkyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkyl group, a substituted or unsubstituted C_3 - C_{10} cycloalkenyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkenyl group, a substituted or unsubstituted C_6 - C_{60} aryl group, a substituted or unsubstituted C_6 - C_{60} aryloxy group, a substituted or unsubstituted C_6 - C_{60} arylthio group, a substituted or unsubstituted C_1 - C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group.

In some embodiments, in Formulae 201 and 202,

L_{201} to L_{205} may each independently be selected from the group consisting of:

a phenylene group, a naphthylene group, a fluorenylene group, spiro-bifluorenylene group, a benzofluorenylene group, a dibenzofluorenylene group, a phenanthrenylene group, an anthracenylylene group, a pyrenylene group, a chrysenylene group, a pyridinylylene group, a pyrazinylylene group, a pyrimidinylene group, a pyridazinylene group, a quinolinylene group, an isoquinolinylene group, a quinoxalinylylene group, a quinazolinylylene group, a carbazolylene group, and a triazinylene group, and

a phenylene group, a naphthylene group, a fluorenylene group, spiro-bifluorenylene group, a benzofluorenylene group, a dibenzofluorenylene group, a phenanthrenylene group, an anthracenylylene group, a pyrenylene group, a chrysenylene group, a pyridinylylene group, a pyrazinylylene group, a pyrimidinylene group, a pyridazinylene group, a quinolinylene group, an isoquinolinylene group, a quinoxalinylylene group, a quinazolinylylene group, a carbazolylene group, and a triazinylene group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{20} alkyl group, a C_1 - C_{20} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinylyl group, a quinazolinylyl group, a carbazolyl group, and a triazinyl group;

$xa1$ to $xa4$ may each independently be 0, 1, or 2;

$xa5$ may be 1, 2, or 3;

R_{201} to R_{204} may each independently be selected from the group consisting of:

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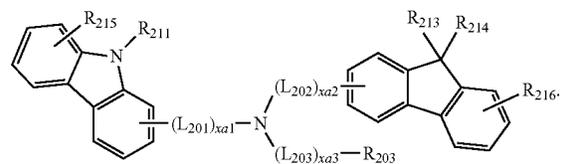
a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinylyl group, a quinazolinylyl group, a carbazolyl group, and a triazinyl group, and

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinylyl group, a quinazolinylyl group, a carbazolyl group, and a triazinyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{20} alkyl group, a C_1 - C_{20} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, an azulenyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinylyl group, a quinazolinylyl group, a carbazolyl group, and a triazinyl group;

but embodiments of the present disclosure are not limited thereto.

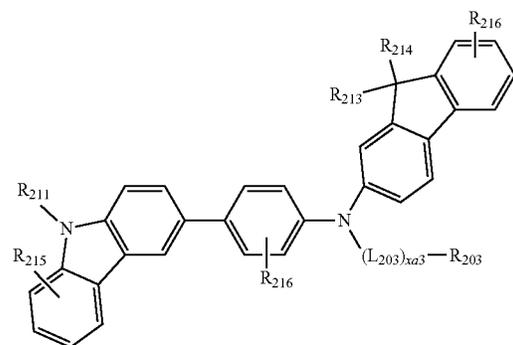
The compound represented by Formula 201 may be represented by Formula 201A:

Formula 201A



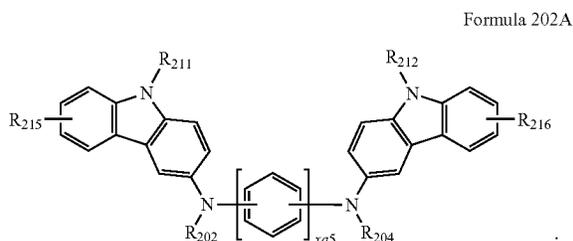
For example, the compound represented by Formula 201 may be represented by Formula 201A-1 below, but is not limited thereto:

Formula 201A-1



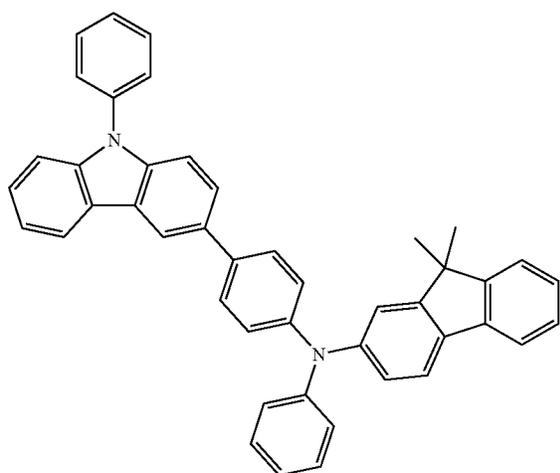
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For example, the compound represented by Formula 202 may be represented by Formula 202A below, but is not limited thereto:



Regarding Formulae 201A, 201A-1, and 202A, L_{201} to L_{203} , $xa1$ to $xa3$, $xa5$, and R_{202} to R_{204} are already described above; descriptions of R_{211} and R_{212} may each independently be understood by referring to the description of R_{203} ; and R_{213} to R_{216} may each independently be selected from hydrogen, deuterium, $-F$, $-Cl$, $-Br$, $-I$, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{60} alkyl group, a C_2 - C_{60} alkenyl group, a C_2 - C_{60} alkynyl group, a C_1 - C_{60} alkoxy group, a C_3 - C_{10} cycloalkyl group, a C_1 - C_{10} heterocycloalkyl group, a C_3 - C_{10} cycloalkenyl group, a C_1 - C_{10} heterocycloalkenyl group, a C_6 - C_{60} aryl group, a C_6 - C_{60} aryloxy group, a C_6 - C_{60} arylthio group, a C_1 - C_{60} heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group.

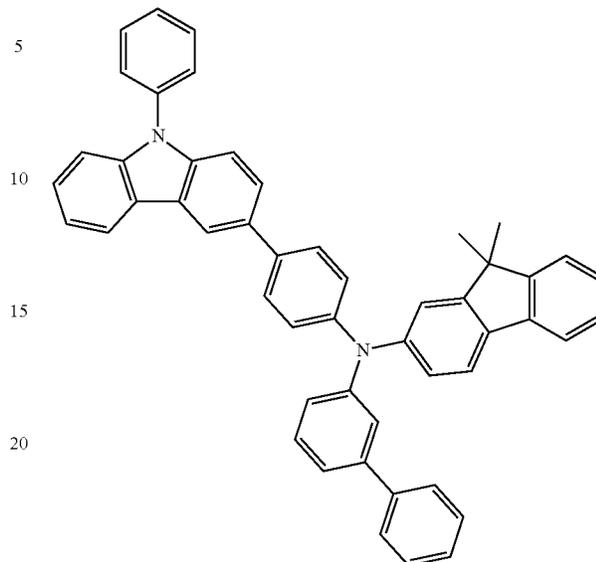
The compound represented by Formula 201, and the compound represented by Formula 202 may each independently include any of compounds HT1 to HT20 illustrated below, but are not limited thereto.



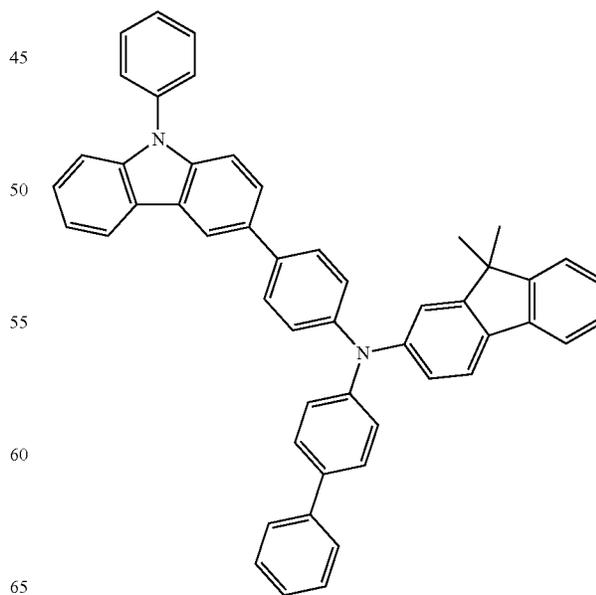
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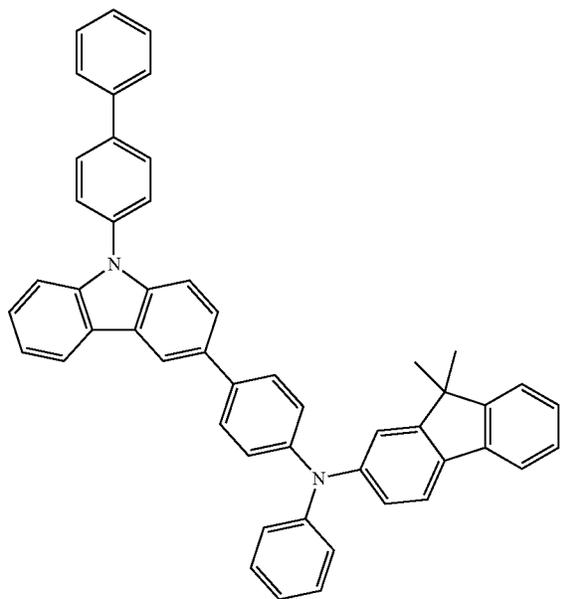
HT2



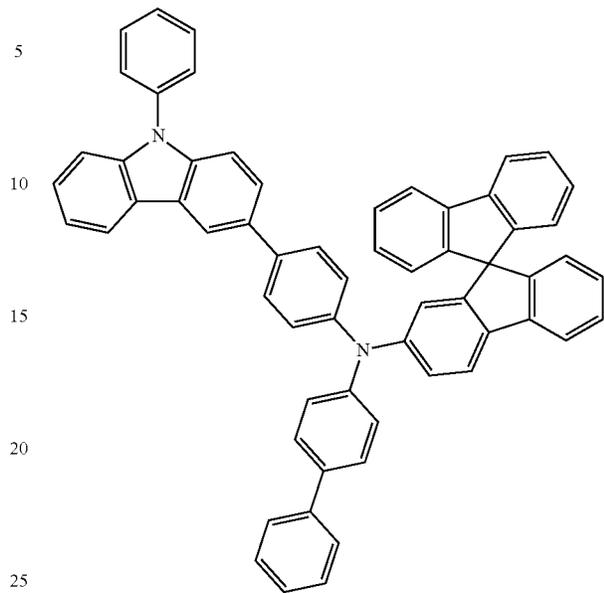
HT3



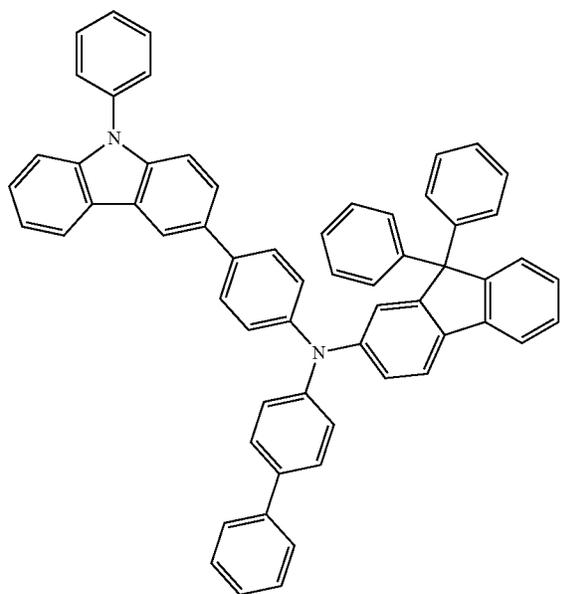
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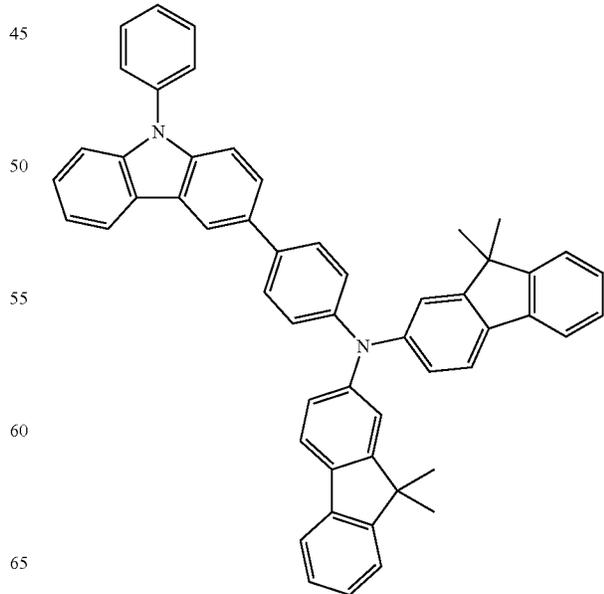
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HT5

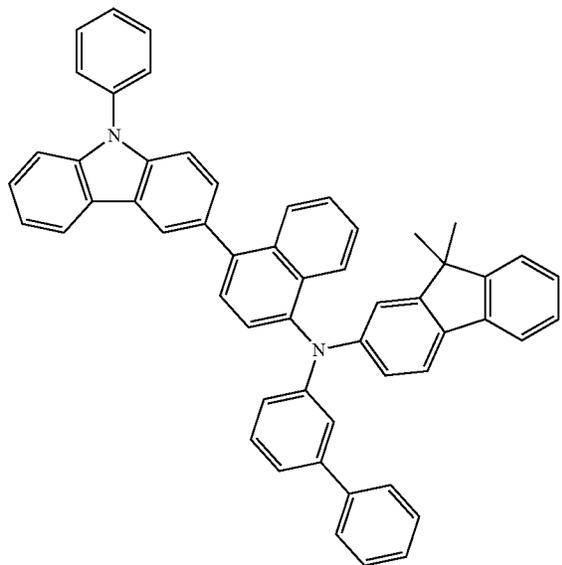


HT7



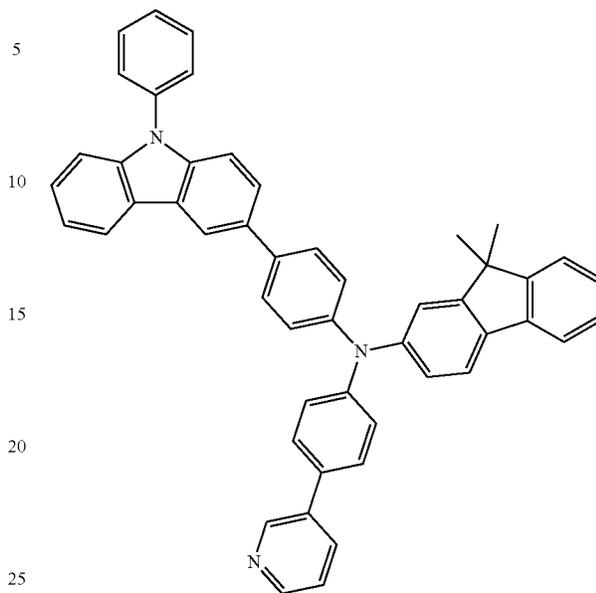
79
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HT8



80
-continued

HT10



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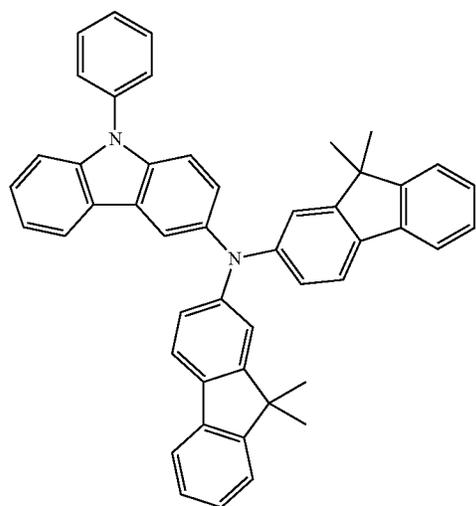
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HT11

HT9



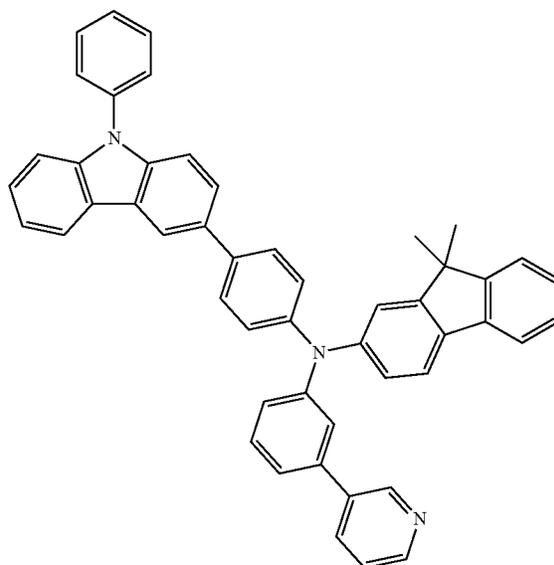
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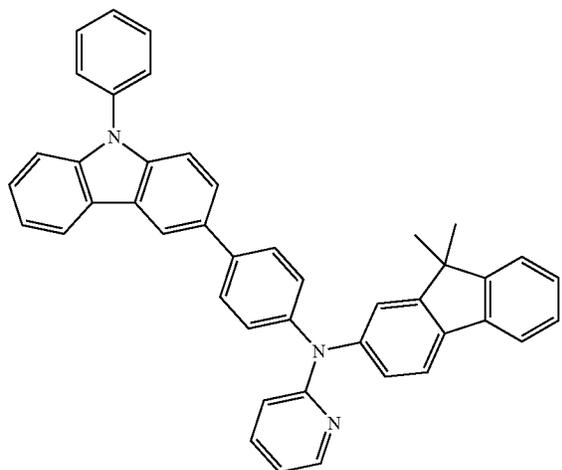
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81

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HT12

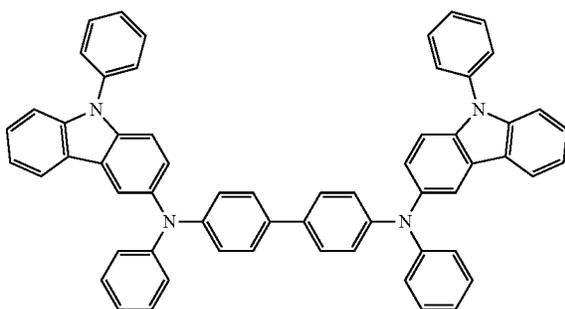


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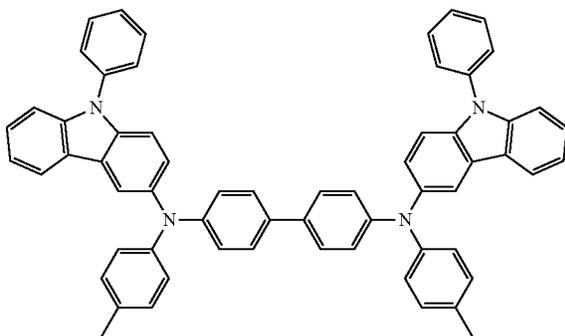
HT13



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HT14

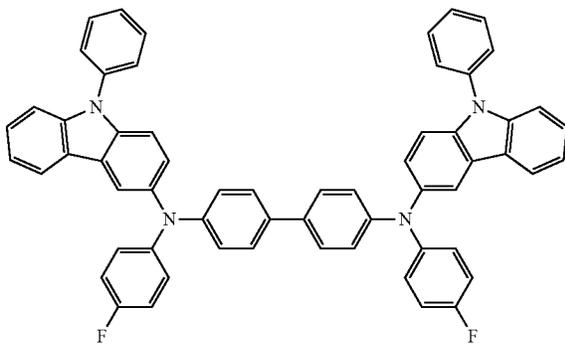


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HT15



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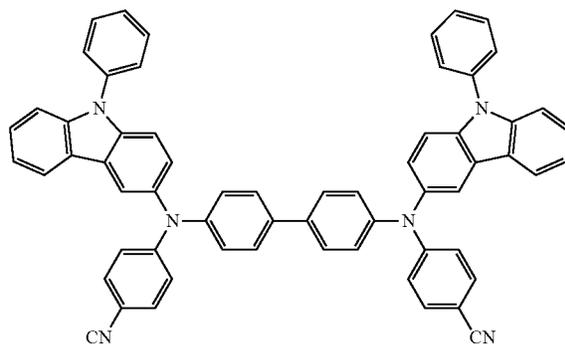
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HT16

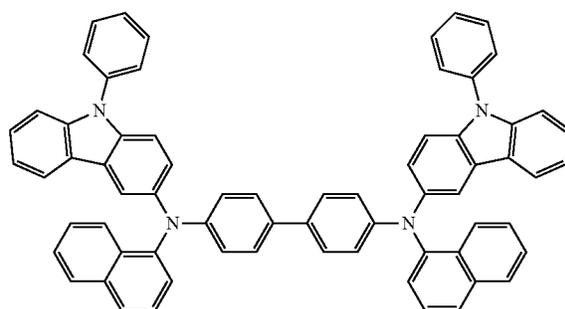


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HT17

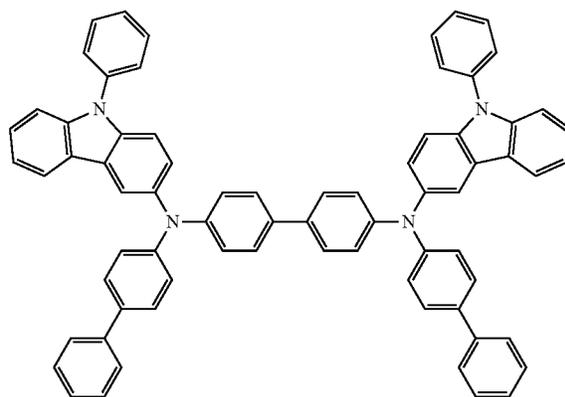


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HT18

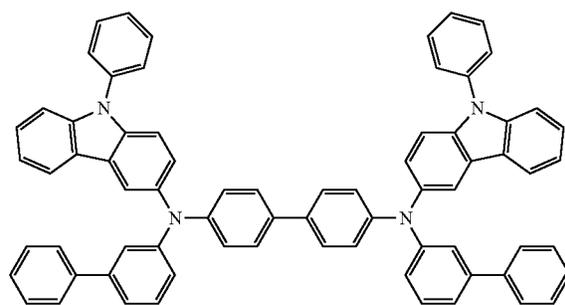


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HT19



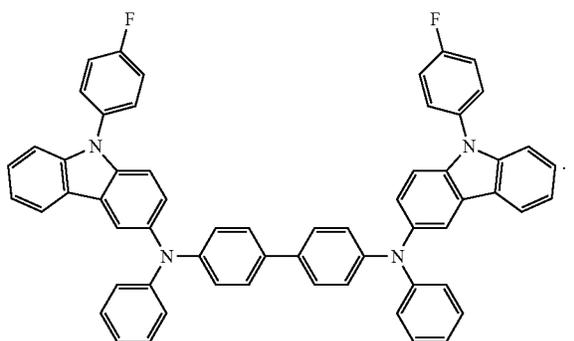
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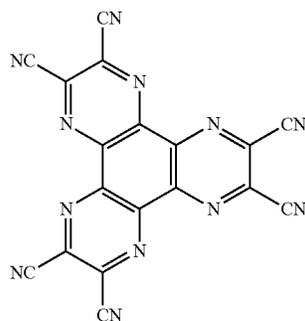
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A thickness of the hole transport region may be in a range of about 100 Å to about 10,000 Å, for example, about 100 Å to about 1,000 Å. When the hole transport region includes at least one selected from a hole injection layer and a hole transport layer, a thickness of the hole injection layer may be in a range of about 100 Å to about 10,000 Å, for example, about 100 Å to about 1,000 Å, and a thickness of the hole transport layer may be in a range of about 50 Å to about 2,000 Å, for example, about 100 Å to about 1,500 Å. When the thicknesses of the hole transport region, the hole injection layer, and the hole transport layer are within any of these ranges, satisfactory (or suitable) hole transporting characteristics may be obtained without a substantial increase in driving voltage.

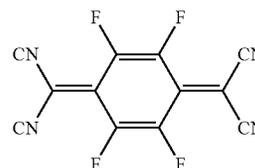
The hole transport region may further include, in addition to the materials described above, a charge-generation material for the improvement of conductive properties. The charge-generation material may be homogeneously or unhomogeneously dispersed in the hole transport region.

The charge-generation material may be, for example, a p-dopant. The p-dopant may be one selected from a quinone derivative, a metal oxide, and a cyano group-containing compound, but embodiments are not limited thereto. Non-limiting examples of the p-dopant include quinone derivatives (such as tetracyanoquinonodimethane (TCNQ) and/or 2,3,5,6-tetrafluoro-tetracyano-1,4-benzoquinonodimethane (F4-TCNQ)), metal oxides (such as tungsten oxide and/or molybdenum oxide), and Compound HT-D1 illustrated below.



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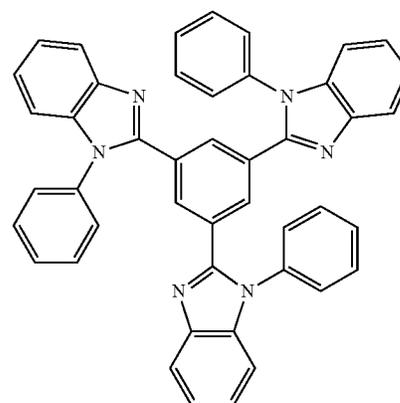
The hole transport region may further include, in addition to the hole injection layer and/or the hole transport layer, at least one selected from a buffer layer and an electron blocking layer. Since the buffer layer may compensate for an optical resonance distance according to a wavelength of light emitted from the emission layer, light-emission efficiency of the formed organic light-emitting device may be improved. For use as a material included in the buffer layer, any of materials that are to be included in the hole transport region may be used. The electron blocking layer may function to prevent or reduce the injection of electrons from the electron transport region.

An emission layer may be formed on the first electrode **110** or the hole transport region by using one or more suitable methods such as vacuum deposition, spin coating, casting, a LB method, ink-jet printing, laser-printing, and/or laser-induced thermal imaging. When an emission layer is formed by vacuum deposition and/or spin coating, deposition and coating conditions for the emission layer may be the same as (or similar to) those for the hole injection layer.

When the organic light-emitting device **10** is a full color organic light-emitting device, the emission layer may be patterned into a red emission layer, a green emission layer, and/or a blue emission layer, according to a sub pixel. In some embodiments, the emission layer may have a stacked structure including a red emission layer, a green emission layer, and a blue emission layer, or may include a red-light emission material, a green-light emission material, and a blue-light emission material, which are mixed with each other in a single layer, to emit white light.

The emission layer may include a host and a dopant.

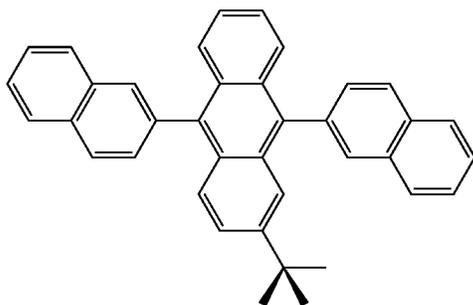
The host may include at least one selected from TPBi, TBADN, ADN (herein also referred to as "DNA"), CBP, CDBP, and TCP:



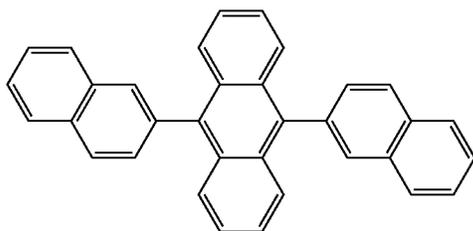
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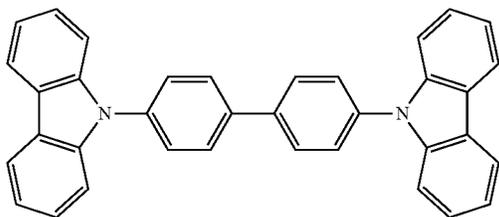
TBADN



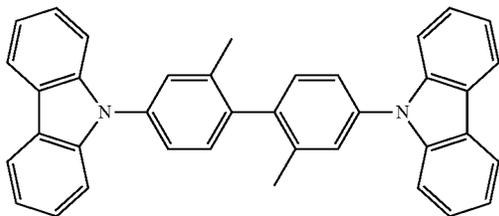
ADN



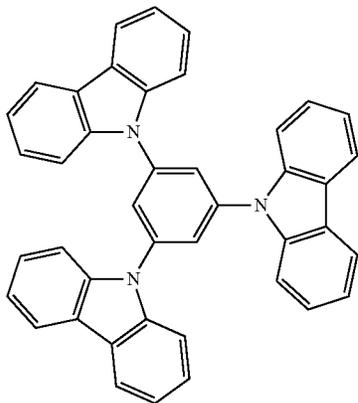
CDBP



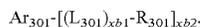
CDBP



TCP



In some embodiments, the host may include a compound represented by Formula 301 below.



Formula 301

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In Formula 301,

Ar₃₀₁ may be selected from the group consisting of:

a naphthalene, a heptalene, a fluorene, a spiro-fluorene, a benzofluorene, a dibenzofluorene, a phenalene, a phenanthrene, an anthracene, a fluoranthene, a triphenylene, a pyrene, a chrysene, a naphthacene, a picene, a perylene, a pentaphene, and an indenoanthracene, and

a naphthalene group, a heptalene group, a fluorene group, a spiro-fluorene group, a benzofluorene group, a dibenzofluorene group, a phenalene group, a phenanthrene group, an anthracene group, a fluoranthene group, a triphenylene group, a pyrene group, a chrysene group, a naphthacene group, a picene group, a perylene group, a pentaphene group, and an indenoanthracene group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I,

a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, and —Si(Q₃₀₁)(Q₃₀₂)(Q₃₀₃) (wherein Q₃₀₁ to Q₃₀₃ may each independently be selected from hydrogen, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₆-C₆₀ aryl group, and a C₁-C₆₀ heteroaryl group);

description of L₃₀₁ may be the same as the description provided in connection with L₁;

R₃₀₁ may be selected from the group consisting of:a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group,

a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group, and

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine

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group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazoliny

group, a carbazolyl group, and a triazinyl group;

xb1 may be selected from 0, 1, 2, and 3; and

xb2 may be selected from 1, 2, 3, and 4.

For example, in Formula 301,

L₃₀₁ may be selected from the group consisting of:

a phenylene group, a naphthylene group, a fluorenylene group, a spiro-fluorenylene group, a benzofluorenylene group, a dibenzofluorenylene group, a phenanthrenylene group, an anthracenylylene group, a pyrenylene group, and a chrysenylene group, and

a phenylene group, a naphthylene group, a fluorenylene group, a spiro-fluorenylene group, a benzofluorenylene group, a dibenzofluorenylene group, a phenanthrenylene group, an anthracenylylene group, a pyrenylene group, and a chrysenylene group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, and a chrysenyl group;

R₃₀₁ may be selected from the group consisting of:

a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group,

a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, and a chrysenyl group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, and a chrysenyl group, and

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, and a chrysenyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzo-

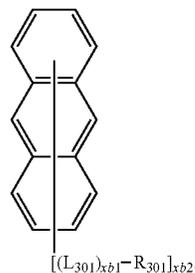
88

fluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, and a chrysenyl group,

but embodiments of the present disclosure are not limited thereto.

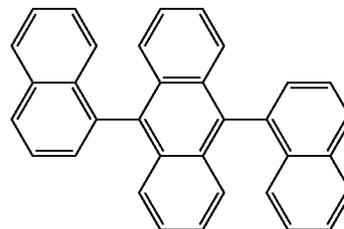
For example, the host may include a compound represented by Formula 301A below:

Formula 301A

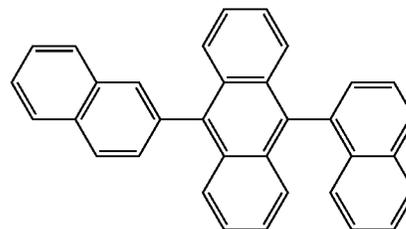


Descriptions of substituents of Formula 301A may be understood by referring to the descriptions thereof provided herein.

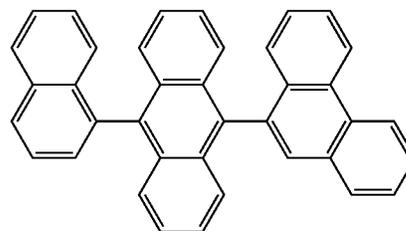
The compound represented by Formula 301 may include at least one of Compounds H1 to H42, but is not limited thereto:



H1



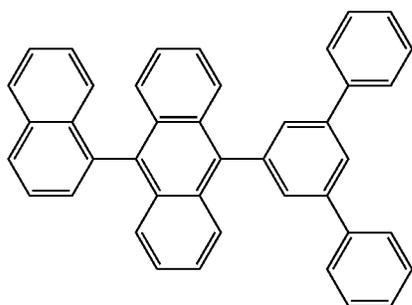
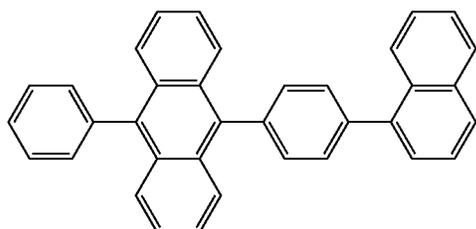
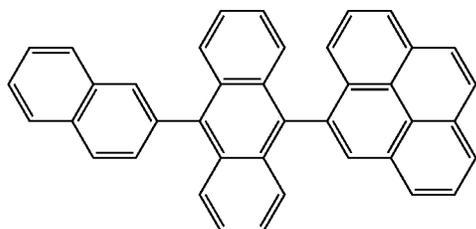
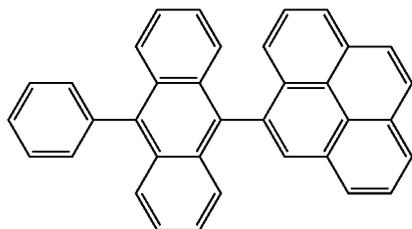
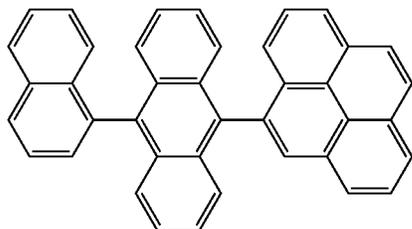
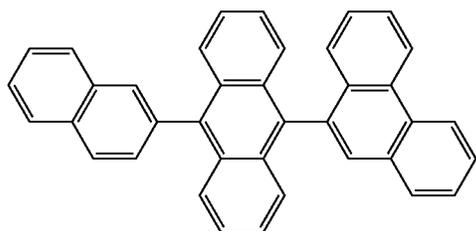
H2



H3

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H4

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H5

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H6

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H7

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H8

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H9

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H10

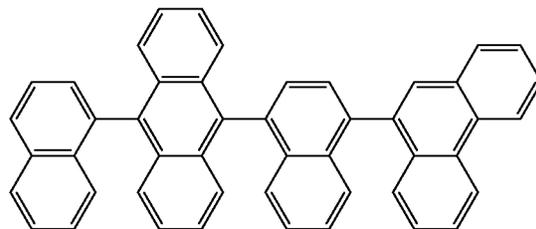
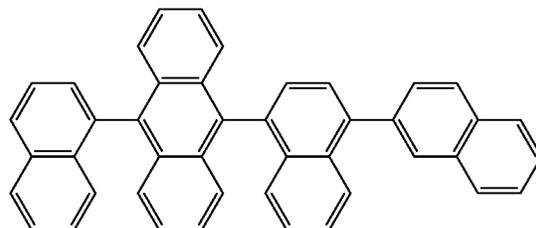
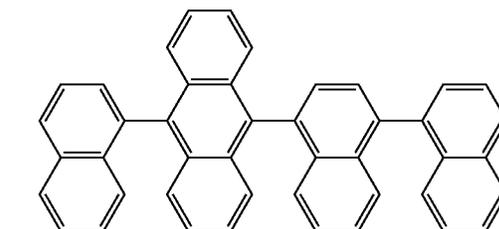
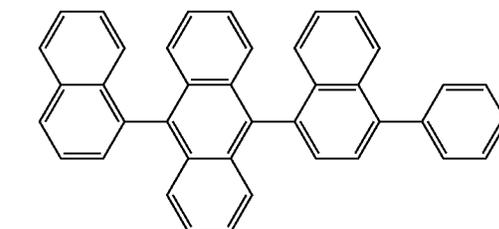
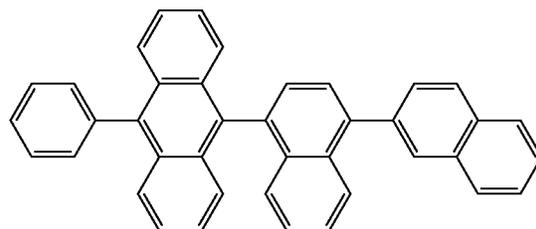
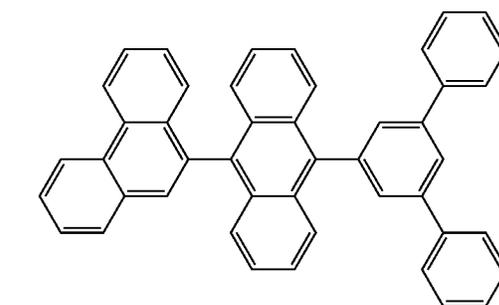
H11

H12

H13

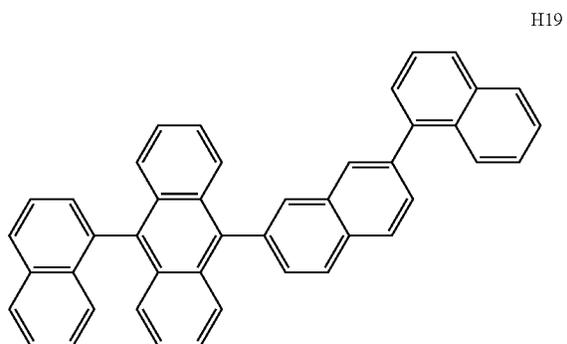
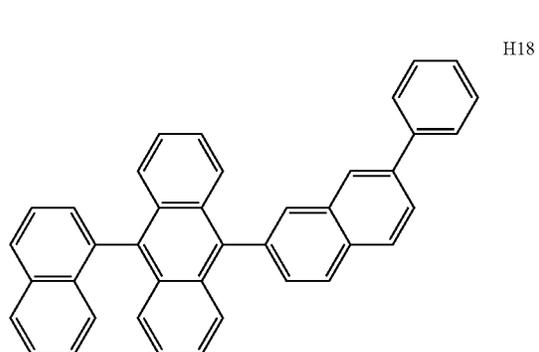
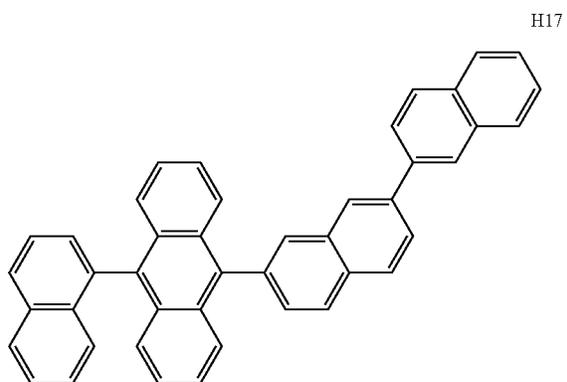
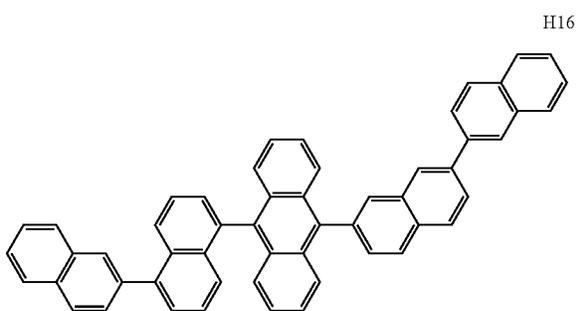
H14

H15



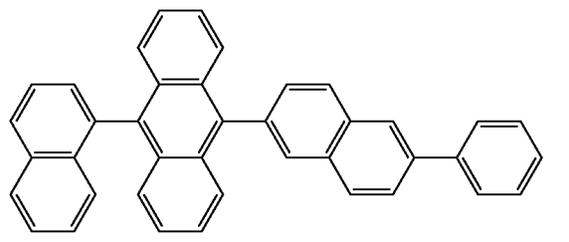
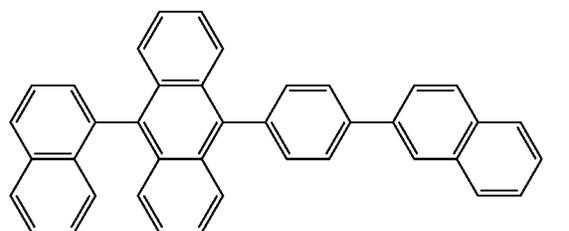
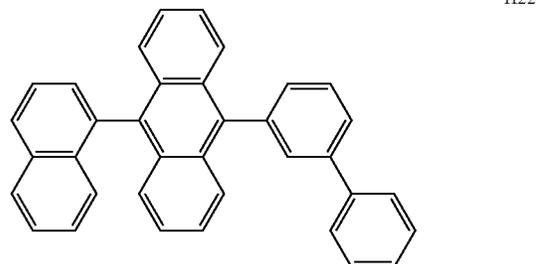
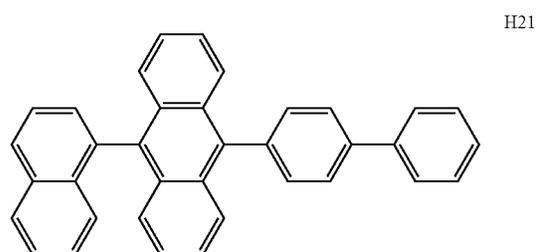
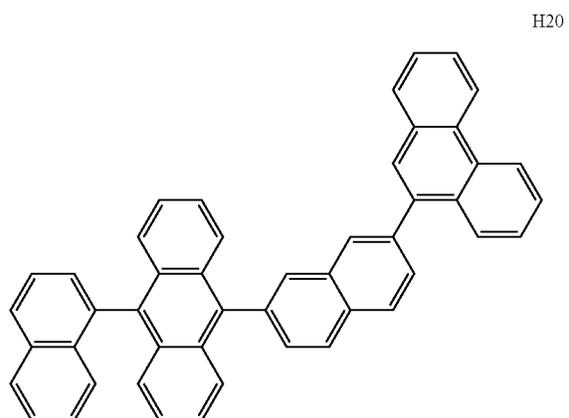
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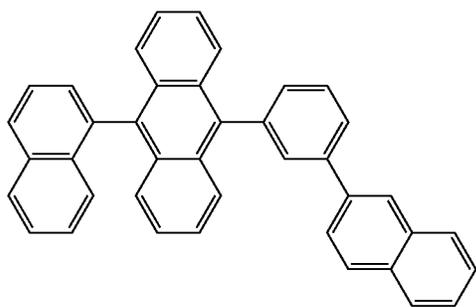
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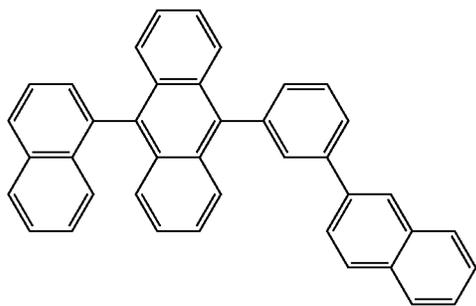
H25

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H26

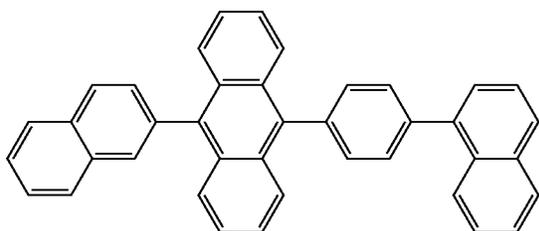
15



H27

20

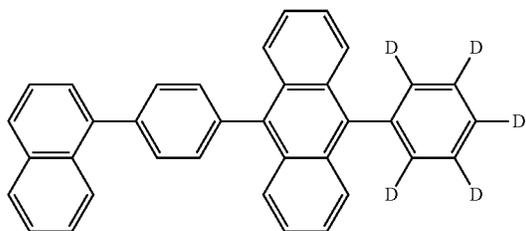
25



H28

30

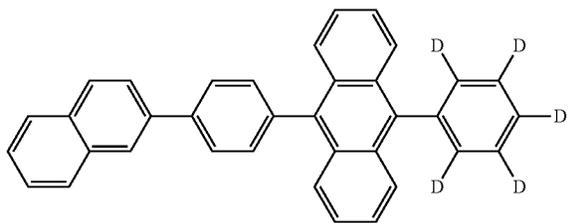
35



H29

40

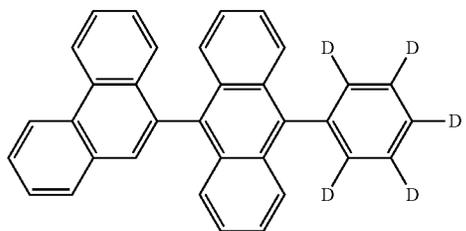
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H30

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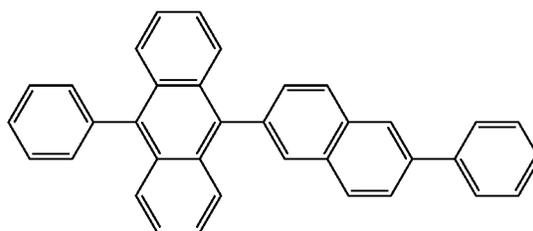


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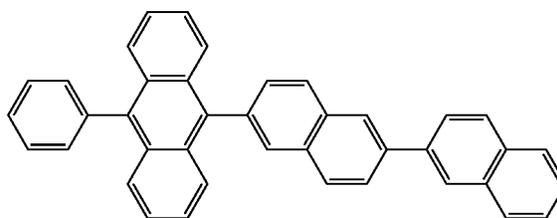
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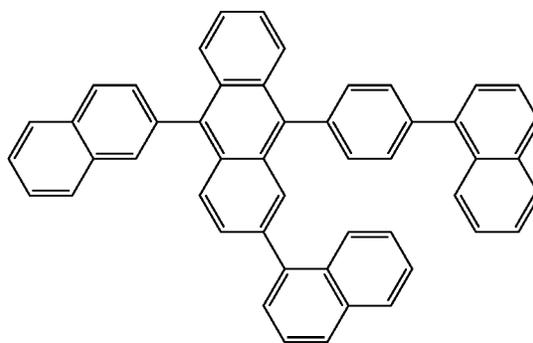
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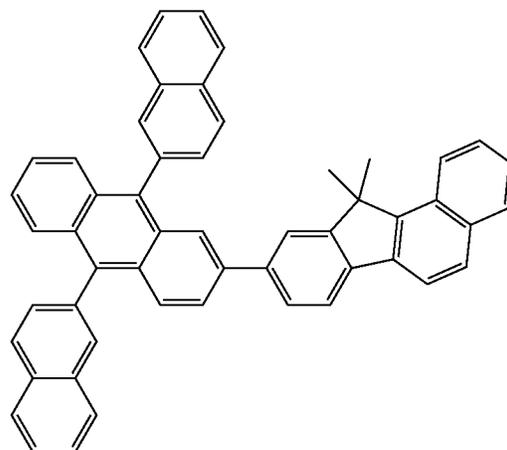
H31



H32



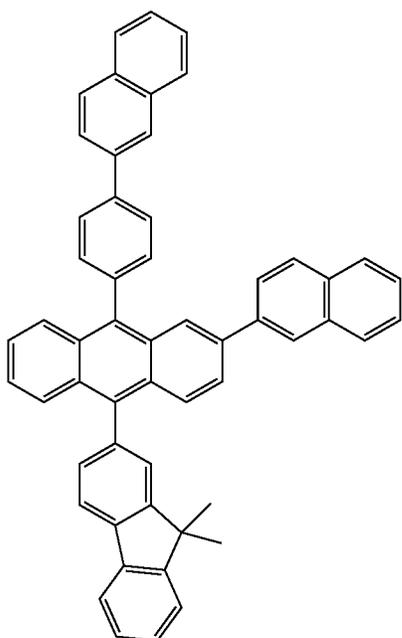
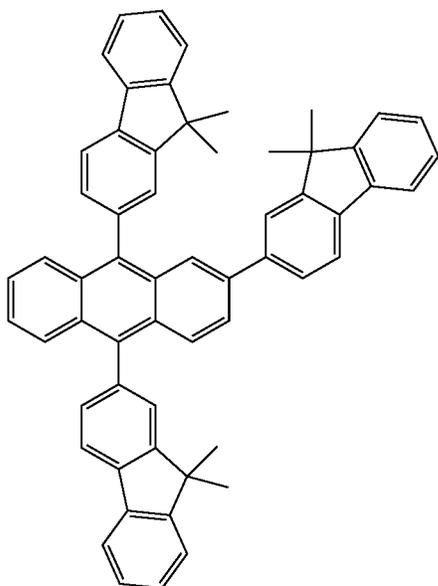
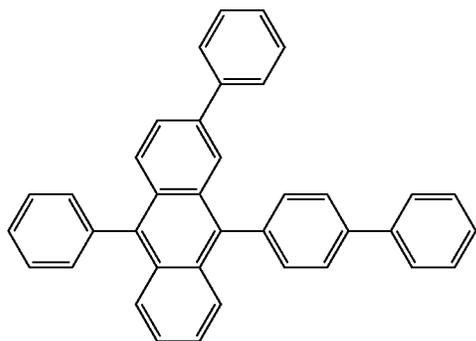
H33



H34

95

-continued



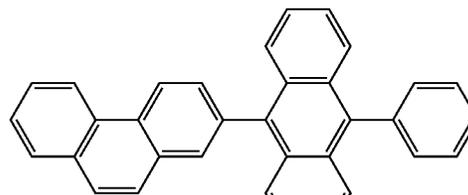
96

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H35

H38

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H36

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H37

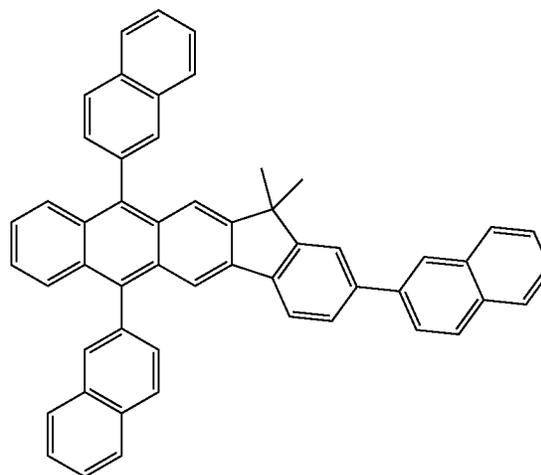
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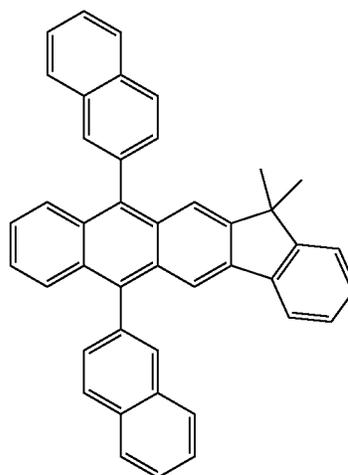
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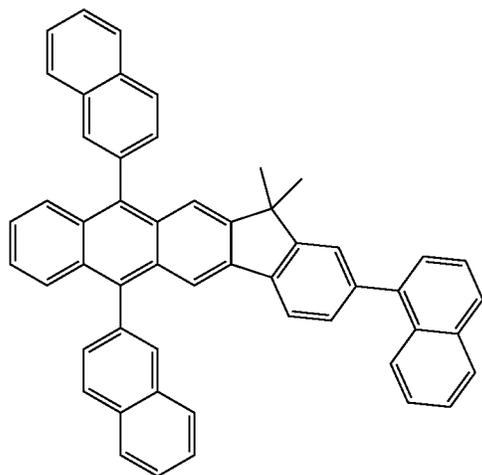


H39



H40

97
-continued



H41

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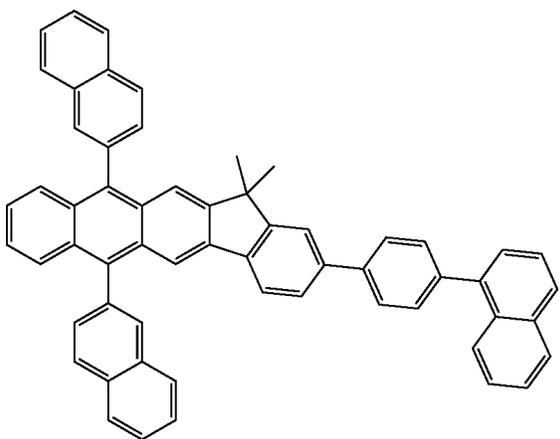
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H42



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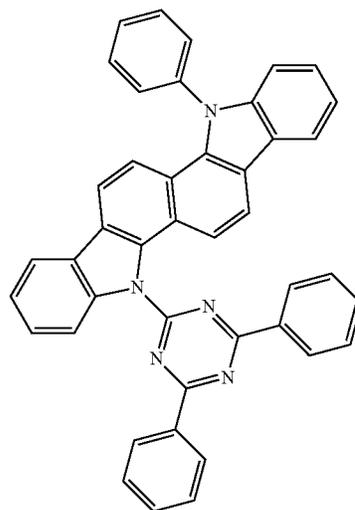
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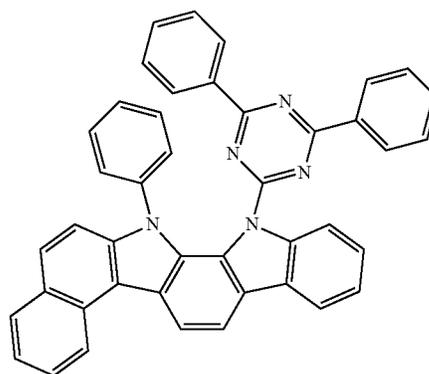
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In some embodiments, the host may include at least one of Compounds H43 to H49 below, but is not limited thereto:

98
-continued



H44



H45

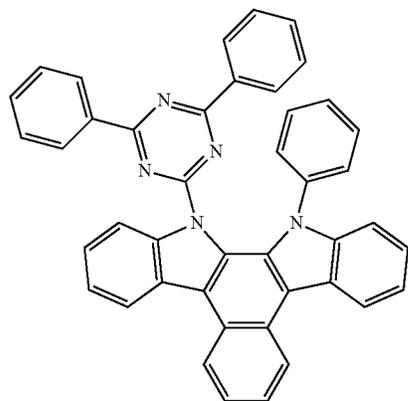
H43

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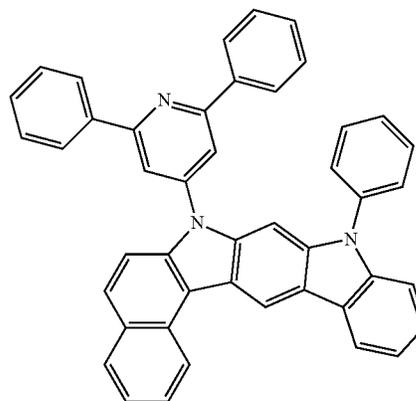
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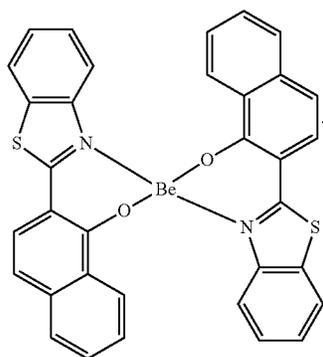
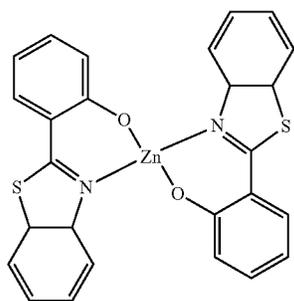
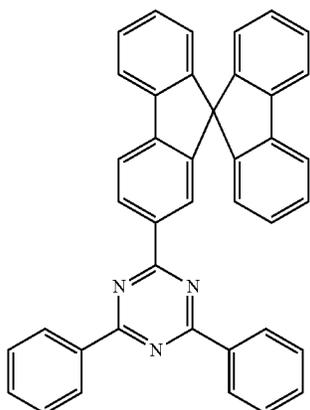


H46



99

-continued



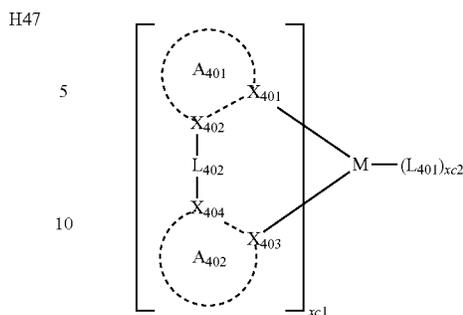
The dopant may include at least one selected from a fluorescent dopant and a phosphorescent dopant.

For example, the phosphorescent dopant may include an organometallic compound including one selected from iridium (Ir), platinum (Pt), osmium (Os), titanium (Ti), zirconium (Zr), hafnium (Hf), europium (Eu), terbium (Tb), thulium (Tm), rhodium (Rh), and copper (Cu).

In some embodiments, the phosphorescent dopant may include an organometallic complex represented by Formula 401 below:

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Formula 401



In Formula 401,

M may be selected from iridium (Ir), platinum (Pt), osmium (Os), titanium (Ti), zirconium (Zr), hafnium (Hf), europium (Eu), terbium (Tb), and thulium (Tm);

X₄₀₁ to X₄₀₄ may each independently be nitrogen or carbon;

A₄₀₁ and A₄₀₂ rings may each independently be selected from a substituted or unsubstituted benzene, a substituted or unsubstituted naphthalene, a substituted or unsubstituted fluorene, a substituted or unsubstituted spiro-fluorene, a substituted or unsubstituted indene, a substituted or unsubstituted pyrrole, a substituted or unsubstituted thiophene, a substituted or unsubstituted furan, a substituted or unsubstituted imidazole, a substituted or unsubstituted pyrazole, a substituted or unsubstituted thiazole, a substituted or unsubstituted isothiazole, a substituted or unsubstituted oxazole, a substituted or unsubstituted isoxazole, a substituted or unsubstituted pyridine, a substituted or unsubstituted pyrazine, a substituted or unsubstituted pyrimidine, a substituted or unsubstituted pyridazine, a substituted or unsubstituted quinoline, a substituted or unsubstituted isoquinoline, a substituted or unsubstituted benzoquinoline, a substituted or unsubstituted quinoxaline, a substituted or unsubstituted quinazoline, a substituted or unsubstituted carbazole, a substituted or unsubstituted benzimidazole, a substituted or unsubstituted benzofuran, a substituted or unsubstituted benzothiophene, a substituted or unsubstituted isobenzothiophene, a substituted or unsubstituted benzoxazole, a substituted or unsubstituted isobenzoxazole, a substituted or unsubstituted triazole, a substituted or unsubstituted oxadiazole, a substituted or unsubstituted triazine, a substituted or unsubstituted dibenzofuran, and a substituted or unsubstituted dibenzothiophene; and

at least one substituent of the substituted benzene, substituted naphthalene, substituted fluorene, substituted spiro-fluorene, substituted indene, substituted pyrrole, substituted thiophene, substituted furan, substituted imidazole, substituted pyrazole, substituted thiazole, substituted isothiazole, substituted oxazole, substituted isoxazole, substituted pyridine, substituted pyrazine, substituted pyrimidine, substituted pyridazine, substituted quinoline, substituted isoquinoline, substituted benzoquinoline, substituted quinoxaline, substituted quinazoline, substituted carbazole, substituted benzimidazole, substituted benzofuran, substituted benzothiophene, substituted isobenzothiophene, substituted benzoxazole, substituted isobenzoxazole, substituted triazole, substituted oxadiazole, substituted triazine, substituted dibenzofuran, and substituted dibenzothiophene may be selected from the group consisting of:

deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic

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acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group,

a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —N(Q₄₀₁)(Q₄₀₂), —Si(Q₄₀₃)(Q₄₀₄)(Q₄₀₅), or —B(Q₄₀₆)(Q₄₀₇),

a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic heterocondensed polycyclic group,

a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₁-C₆₀ alkenyl group, a C₁-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —N(Q₄₁₁)(Q₄₁₂), —Si(Q₄₁₃)(Q₄₁₄)(Q₄₁₅), and —B(Q₄₁₆)(Q₄₁₇), and

—N(Q₄₂₁)(Q₄₂₂), —Si(Q₄₂₃)(Q₄₂₄)(Q₄₂₅), and —B(Q₄₂₆)(Q₄₂₇),

where descriptions of Q₄₀₁ to Q₄₀₇, Q₄₁₁ to Q₄₁₇, and Q₄₂₁ to Q₄₂₇ may each independently be understood by referring to the description provided in connection with Q₁;

L₄₀₁ may be an organic ligand;

xc1 may be 1, 2, or 3; and

xc2 may be 0, 1, 2, or 3.

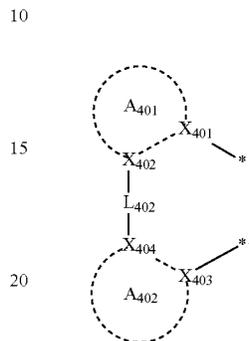
L₄₀₁ may be a monovalent, divalent, or trivalent organic ligand. For example, L₄₀₁ may be selected from a halogen ligand (e.g., Cl and/or F), a diketone ligand (e.g., acetylacetonate, 1,3-diphenyl-1,3-propanedionate, 2,2,6,6-tetramethyl-3,5-heptanedionate, and/or hexafluoroacetonate), a carboxylic acid ligand (e.g., picolinate, dimethyl-3-pyrazolecarboxylate, and/or benzoate), a carbon monoxide ligand, an isonitrile ligand, a cyano ligand, and a phosphorous ligand (e.g., phosphine and/or phosphite), but is not limited thereto.

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When A₄₀₁ in Formula 401 has two or more substituents, the substituents of A₄₀₁ may be linked to form a saturated or unsaturated ring.

When A₄₀₂ in Formula 401 has two or more substituents, the substituents of A₄₀₂ may be linked to form a saturated or unsaturated ring.

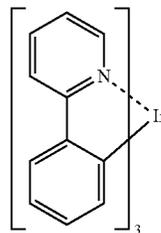
When xc1 in Formula 401 is two or more, a plurality of ligands



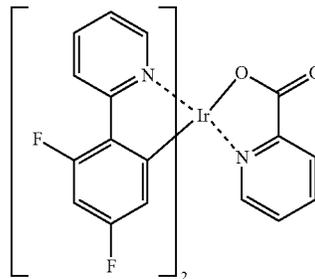
in Formula 401 may be identical to or different from each other. When xc1 in Formula 401 is two or more, A₄₀₁ and A₄₀₂ of one ligand may each independently be connected to respective A₄₀₁ and A₄₀₂ of other neighboring ligands, either directly (e.g., via a bond such as a single bond) or with a linker (e.g., a C₁-C₅ alkylene group, —N(R')— (wherein R' may be a C₁-C₁₀ alkyl group or a C₆-C₂₀ aryl group) and/or —C(=O)—) therebetween.

The phosphorescent dopant may include at least one selected from Compounds PD1 to PD74 below, but is not limited thereto:

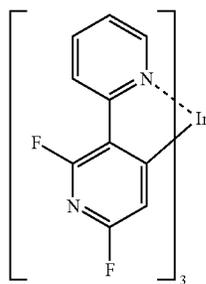
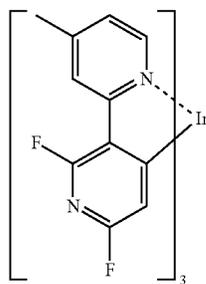
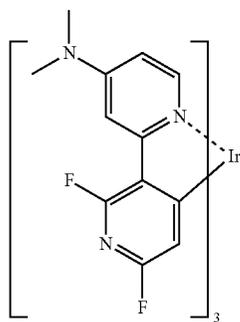
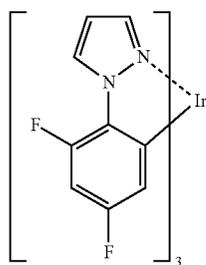
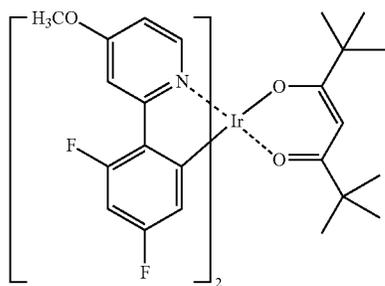
PD1



PD2

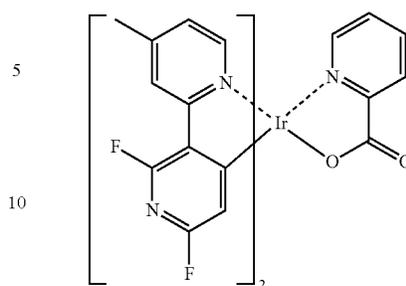


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104
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PD3

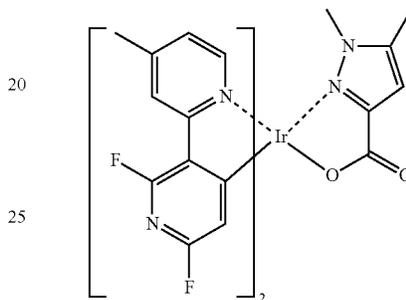


PD8

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PD4

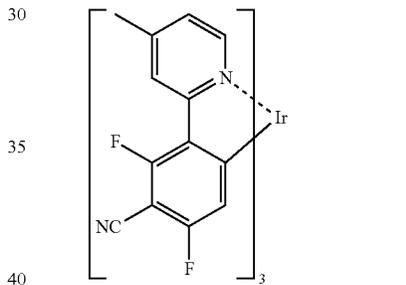


PD9

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PD5

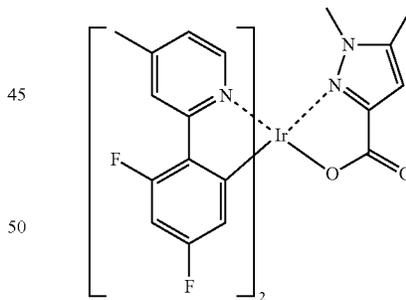


PD10

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PD6

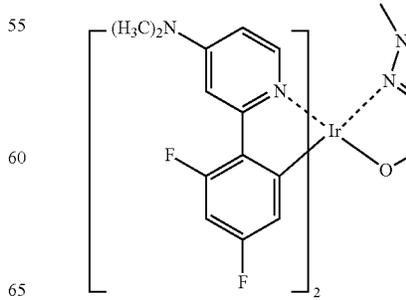


PD11

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PD7



PD12

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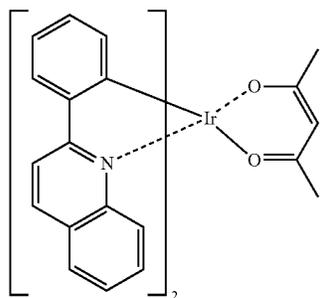
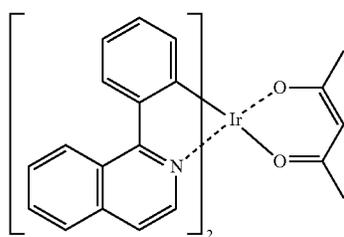
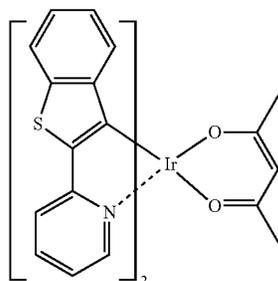
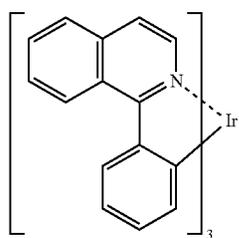
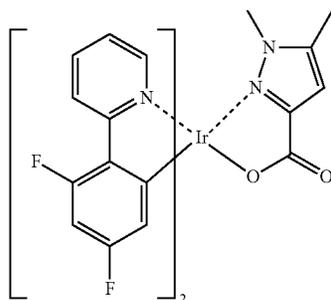
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PD13

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PD16

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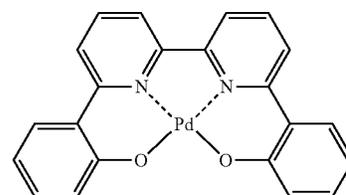
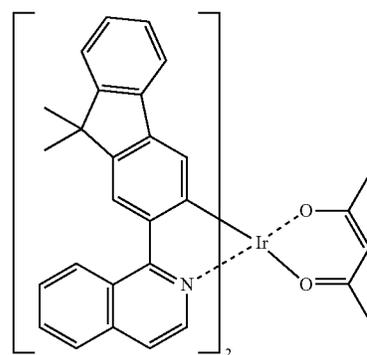
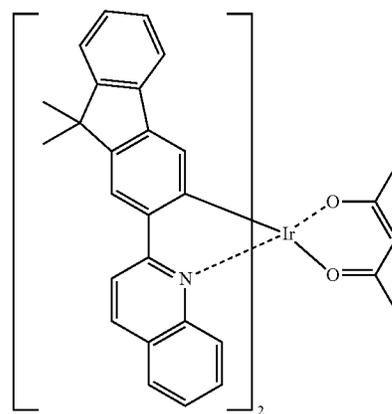
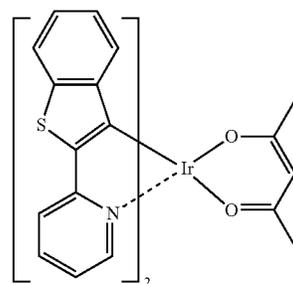
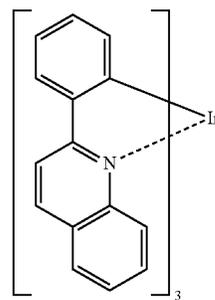
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PD17

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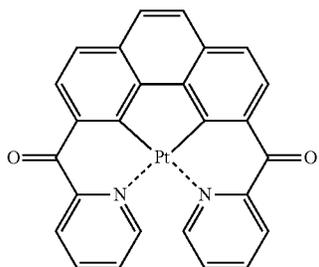
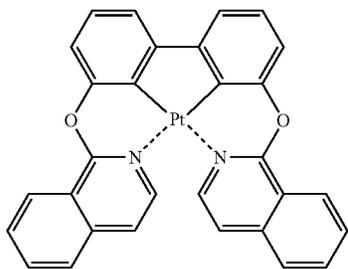
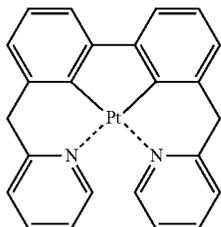
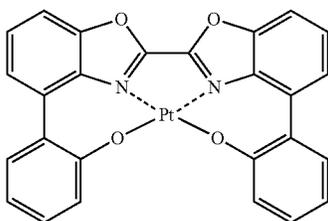
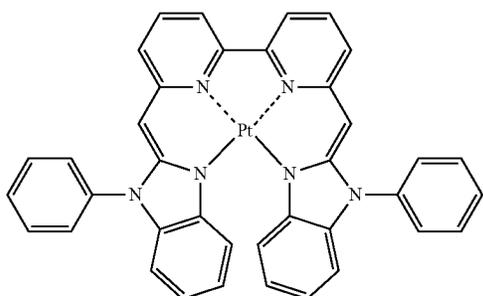
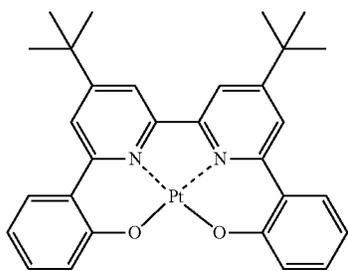
PD20

PD21

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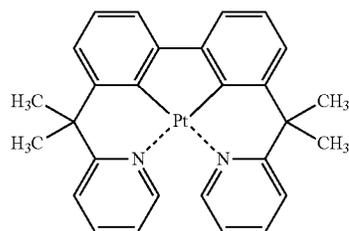


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PD23

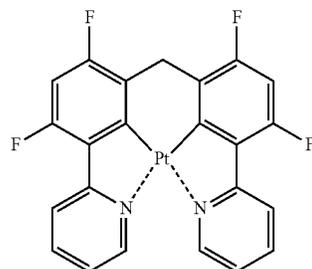
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PD24

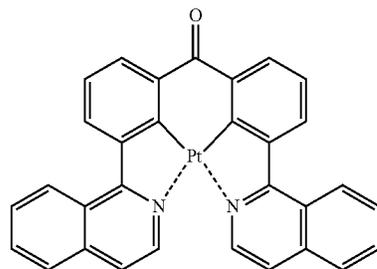
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PD25

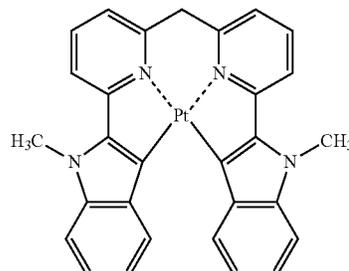
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PD26

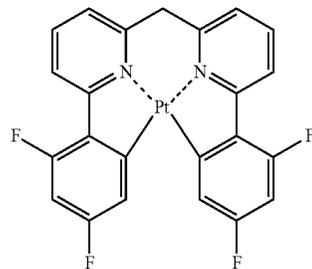
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PD27

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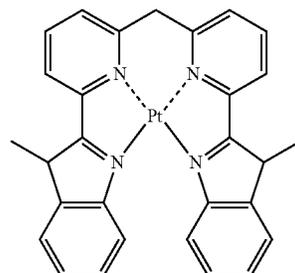


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PD28

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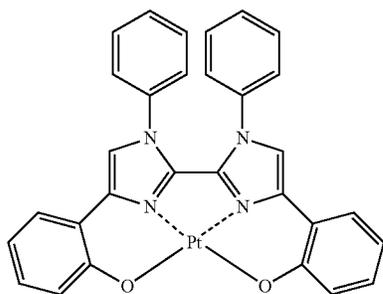
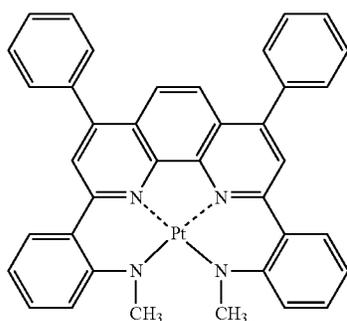
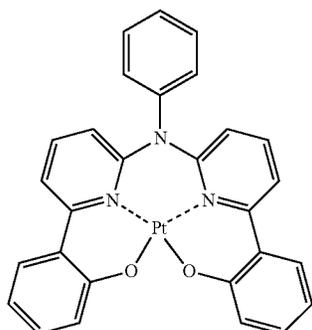
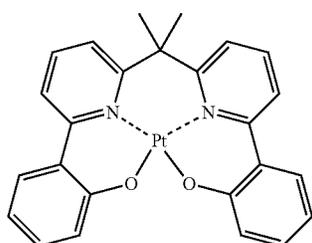
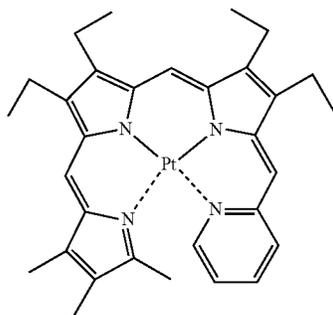
PD31

PD32

PD33

PD34

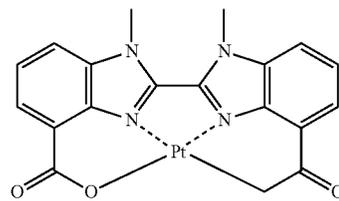
109
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110
-continued

PD35

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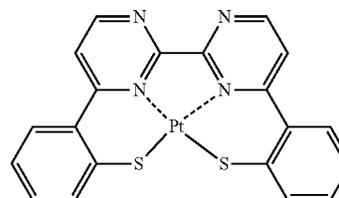


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PD36

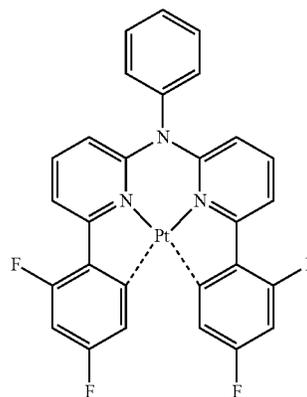
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PD37

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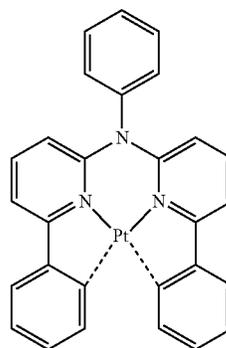


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PD38

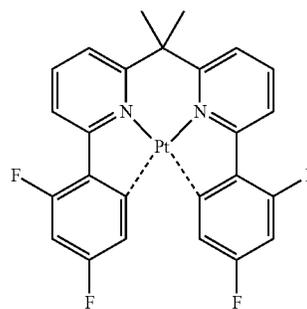
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PD39

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PD40

PD41

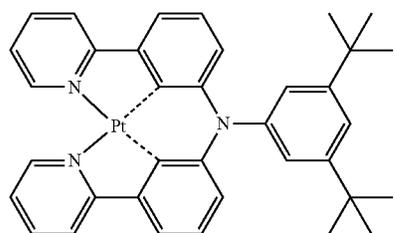
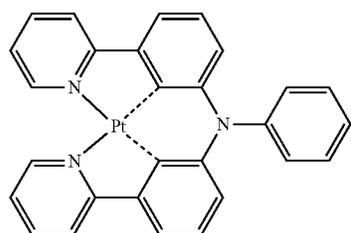
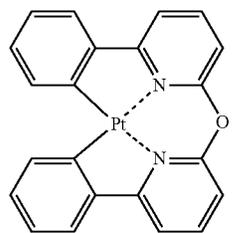
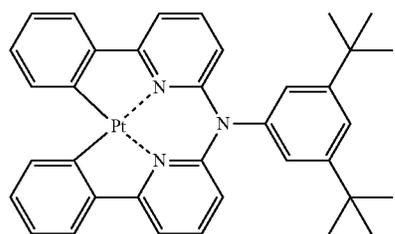
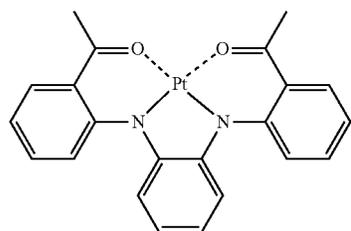
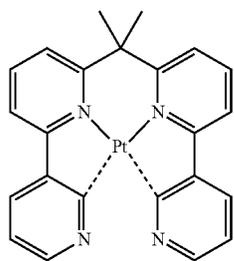
PD42

PD43

PD44

111

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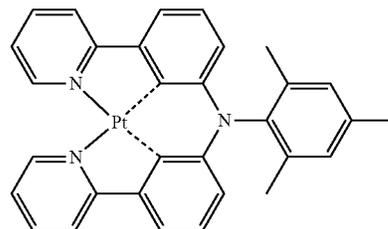
112

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PD45

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PD51

PD46

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PD47

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PD48

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PD49

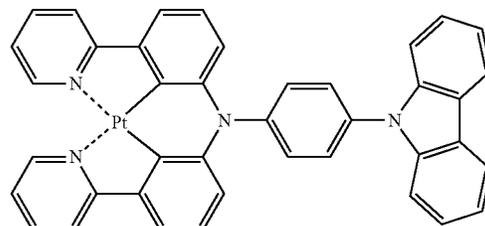
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PD50

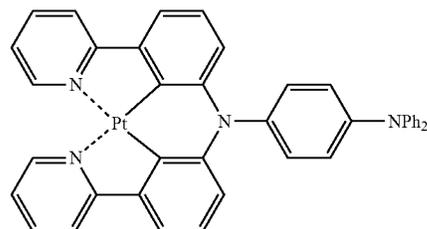
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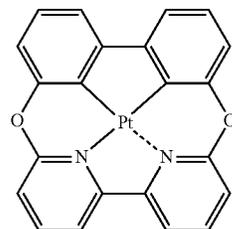


PD52

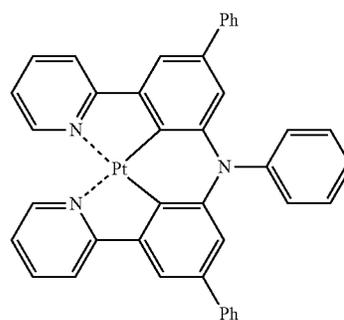
PD53



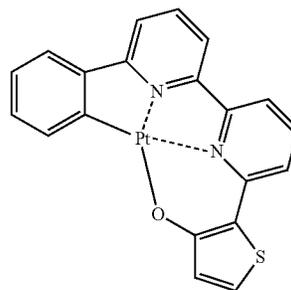
PD54



PD55

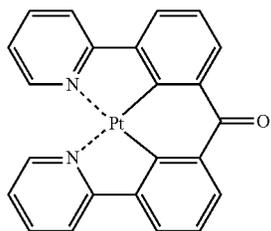


PD56



113

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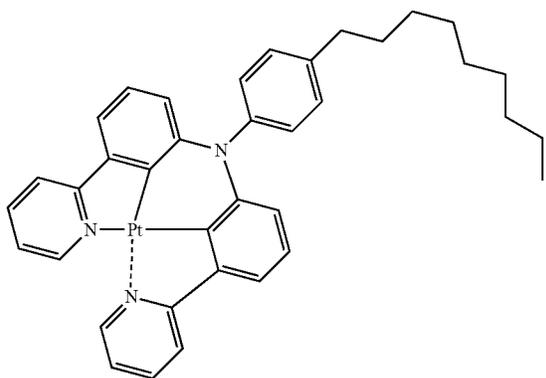
PD57

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PD58

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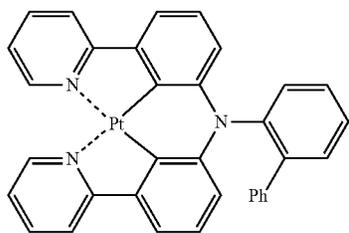
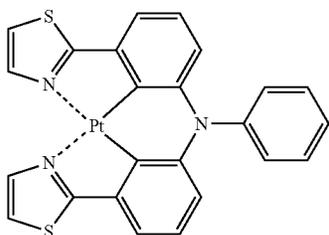
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PD59

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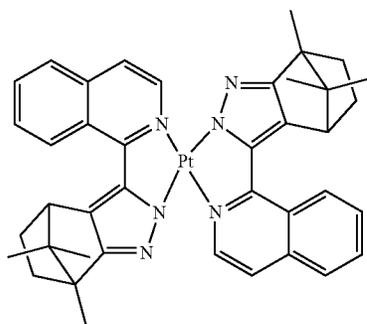
PD60

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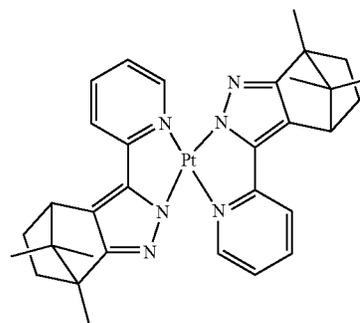
PD61

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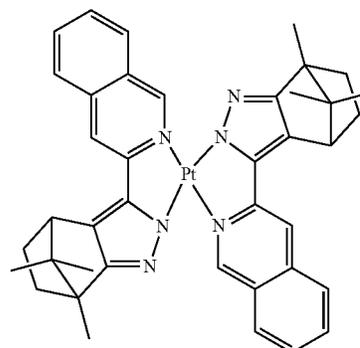


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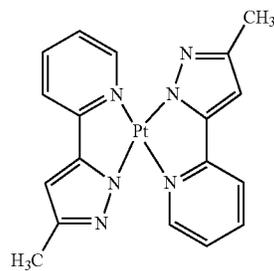
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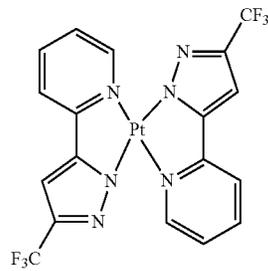
PD62



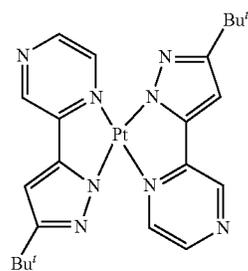
PD63



PD64



PD65



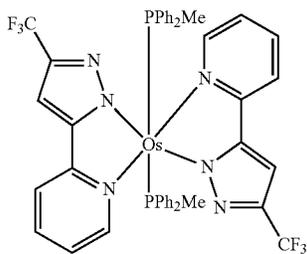
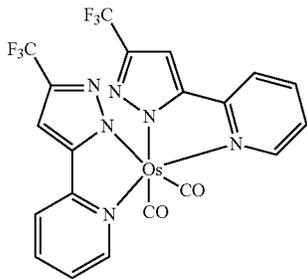
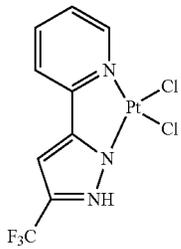
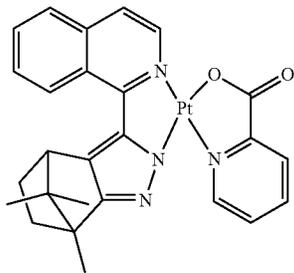
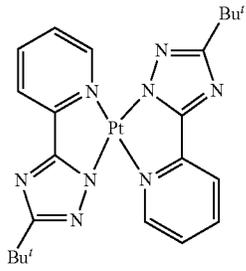
PD66

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115

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116

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PD67

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PD68

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PD69

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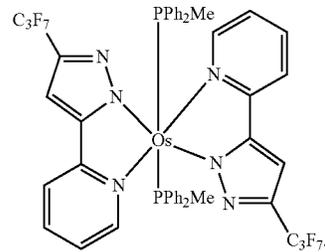
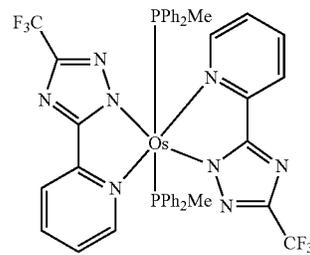
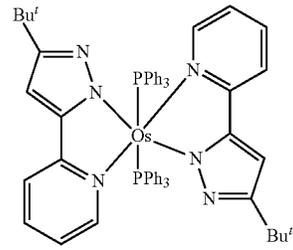
PD70

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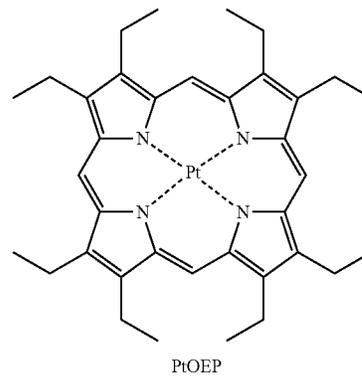
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PD71

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In some embodiments, the phosphorescent dopant may include PtOEP:



The fluorescent dopant may include at least one selected from DPVBi, DPAVBi, TBPe, DCM, DCJTb, Coumarin 6, and C545T.

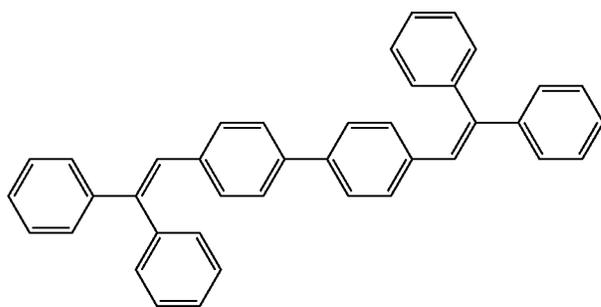
PD72

PD73

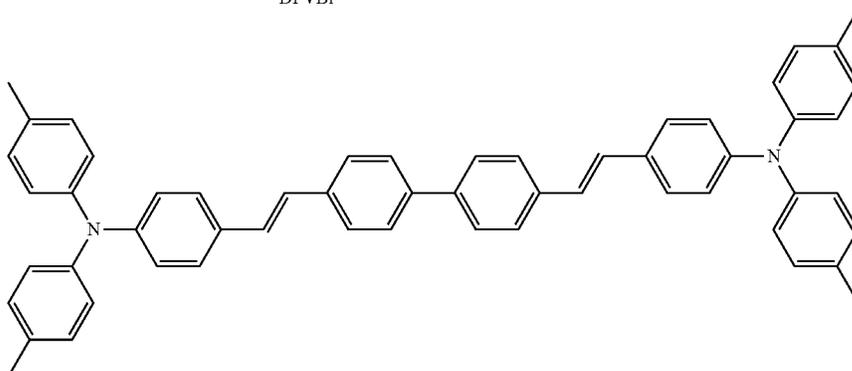
PD74

117

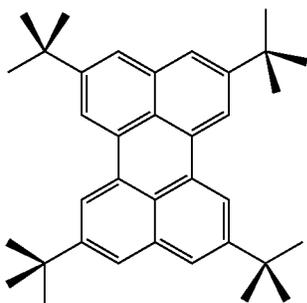
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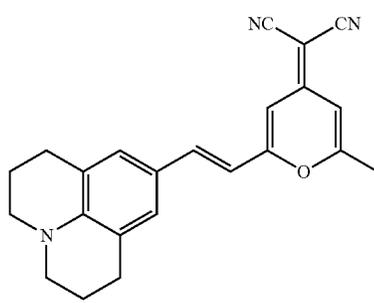
DPVBi



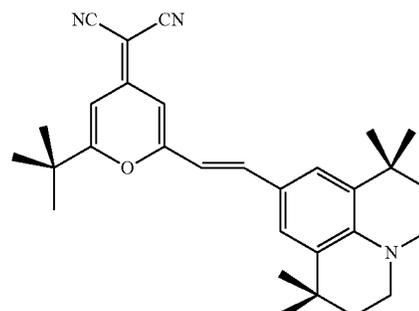
DPAVBi



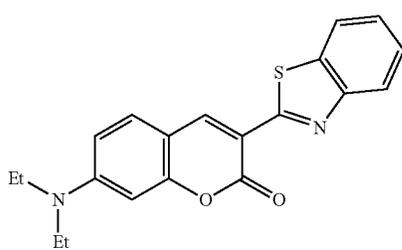
TBPe



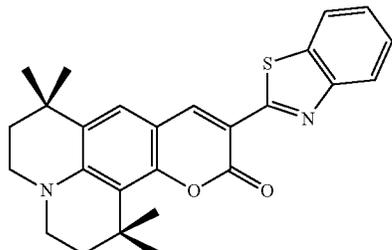
DCM



DCJTb



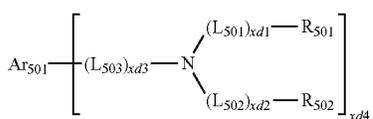
Coumarin 6



C545T

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In some embodiments, the fluorescent dopant may include a compound represented by Formula 501 below.



Formula 501

In Formula 501,

Ar₅₀₁ may be selected from the group consisting of: a naphthalene, a heptalene, a fluorene, a spiro-fluorene, a benzofluorene, a dibenzofluorene, a phenalene, a phenanthrene, an anthracene, a fluoranthene, a triphenylene, a pyrene, a chrysene, a naphthacene, a picene, a perylene, a pentaphene, and an indenoanthracene, and

a naphthalene group, a heptalene group, a fluorene group, a spiro-fluorene group, a benzofluorene group, a dibenzofluorene group, a phenalene group, a phenanthrene group, an anthracene group, a fluoranthene group, a triphenylene group, a pyrene group, a chrysene group, a naphthacene

119

group, a picene group, a perylene group, a pentaphene group, and an indenoanthracene group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, and —Si(Q₅₀₁)(Q₅₀₂)(Q₅₀₃) (wherein Q₅₀₁ to Q₅₀₃ may each independently be selected from hydrogen, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₆-C₆₀ aryl group, and a C₂-C₆₀ heteroaryl group);

descriptions of L₅₀₁ to L₅₀₃ may each independently be understood by referring to the description of L₂₀₁;

R₅₀₁ and R₅₀₂ may each independently be selected from the group consisting of:

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazoliny group, a carbazolyl group, a triazinyl group, a dibenzofuranyl group, and a dibenzothiophenyl group, and

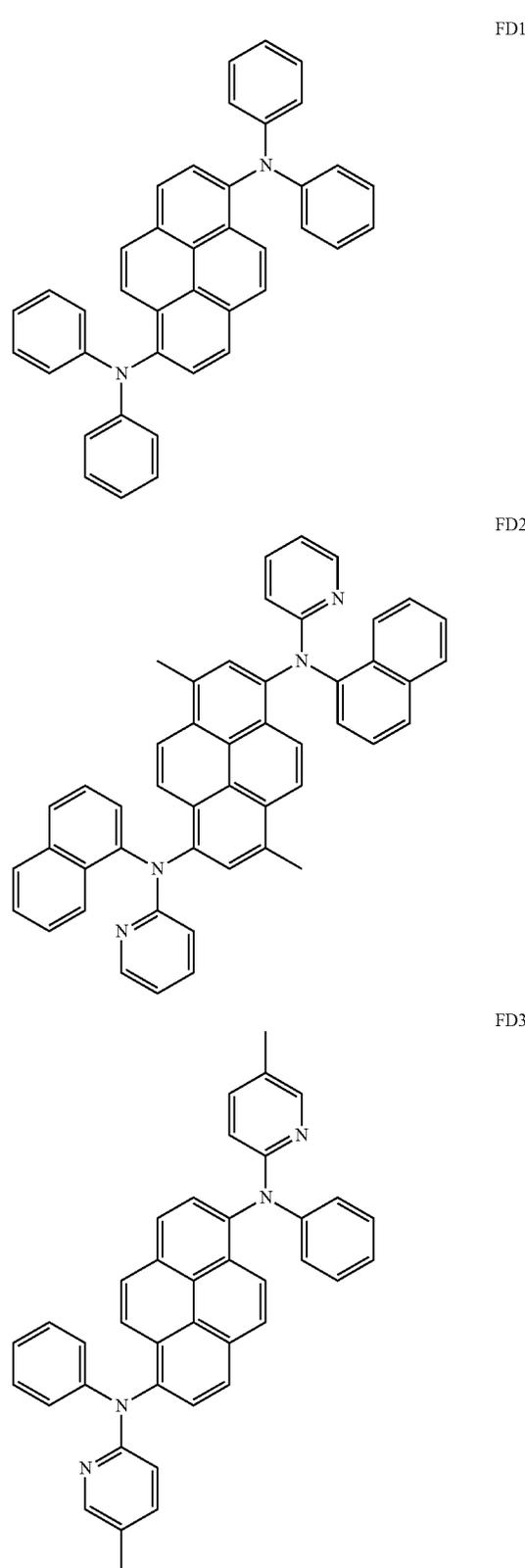
a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazoliny group, a carbazolyl group, a triazinyl group and a dibenzofuranyl group, and a dibenzothiophenyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazoliny group, a carbazolyl group, a triazinyl group, a dibenzofuranyl group, and a dibenzothiophenyl group;

xd1 to xd3 may each independently be selected from 0, 1, 2, and 3; and

xd4 may be selected from 1, 2, 3, and 4.

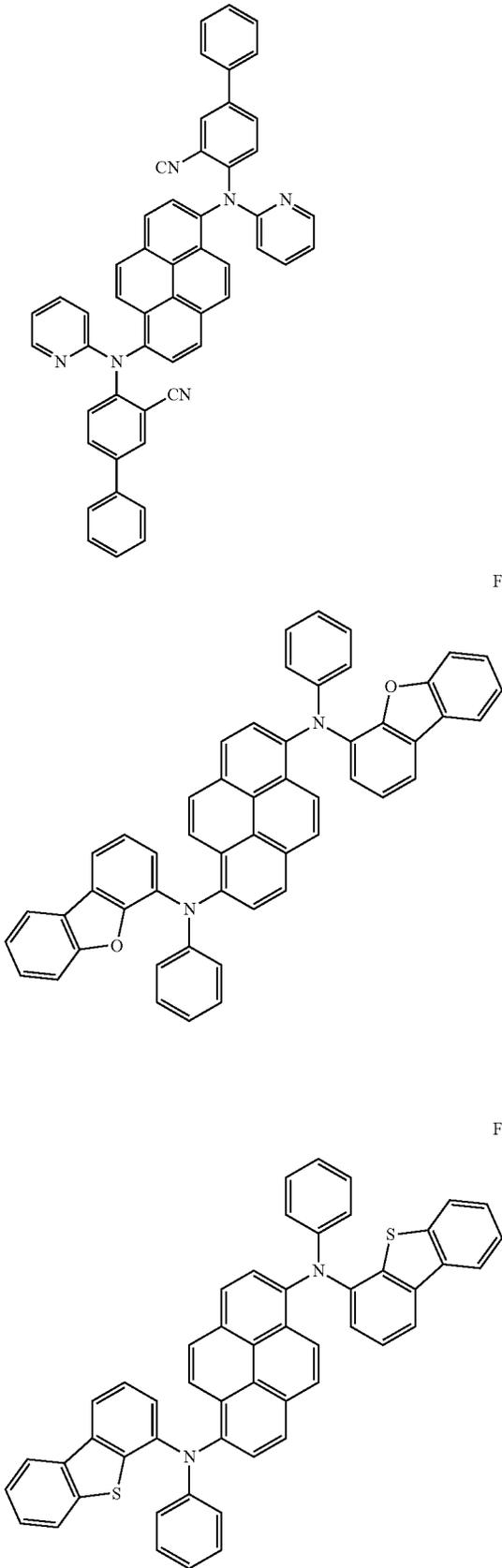
120

The fluorescent dopant may include at least one of Compounds FD1 to FD9:



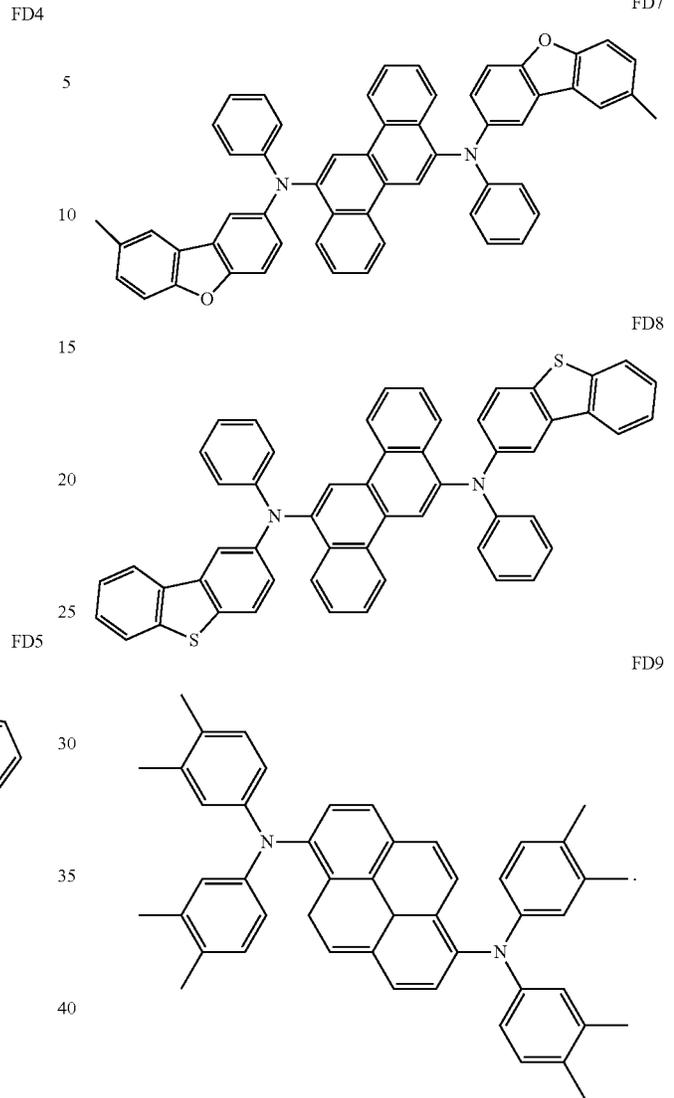
121

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122

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FD4
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FD5
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An amount of the dopant in the emission layer may be, for example, in a range of about 0.01 to about 15 parts by weight based on 100 parts by weight of the host, but is not limited thereto.

FD6
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A thickness of the emission layer may be in a range of about 100 Å to about 1,000 Å, for example, about 200 Å to about 600 Å. When the thickness of the emission layer is within any of these ranges, excellent (or suitable) light-emission characteristics may be obtained without a substantial increase in driving voltage.

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An electron transport region may be disposed on the emission layer.

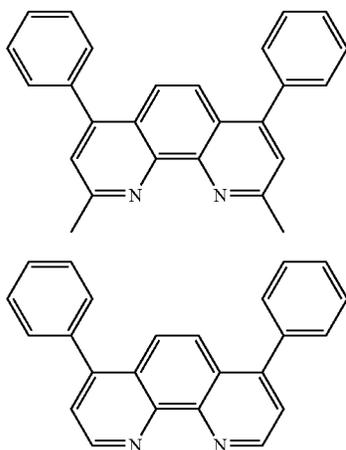
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The electron transport region may include at least one selected from a hole blocking layer, an electron transport layer (ETL), and an electron injection layer, but is not limited thereto.

For example, the electron transport region may have a structure of electron transport layer/electron injection layer or a structure of hole blocking layer/electron transport layer/electron injection layer, wherein the layers of each structure are sequentially stacked in a direction from the emission layer in the stated order, but the structure of the electron transport region is not limited thereto.

According to an embodiment, the organic layer 150 of the organic light-emitting device may include an electron transport region between the emission layer and the second electrode 190.

When the electron transport region includes a hole blocking layer, the hole blocking layer may be formed on the emission layer by using one or more suitable methods such as vacuum deposition, spin coating, casting, a Langmuir-Blodgett (LB) method, ink-jet printing, laser-printing, and/or laser-induced thermal imaging. When the hole blocking layer is formed by vacuum deposition and/or spin coating, deposition and coating conditions for the hole blocking layer may be determined by referring to the deposition and coating conditions for the hole injection layer.

The hole blocking layer may include, for example, at least one selected from BCP and Bphen, but is not limited thereto.



A thickness of the hole blocking layer may be in a range of about 20 Å to about 1,000 Å, for example, about 30 Å to about 300 Å. When the thickness of the hole blocking layer is within any of these ranges, the hole blocking layer may have excellent (or suitable) hole blocking characteristics without a substantial increase in driving voltage.

The electron transport region may include an electron transport layer. The electron transport layer may be formed on the emission layer or the hole blocking layer by using one or more suitable methods such as vacuum deposition, spin coating, casting, a LB method, ink-jet printing, laser-printing, and/or laser-induced thermal imaging. When an electron transport layer is formed by vacuum deposition and/or spin coating, deposition and coating conditions for the electron transport layer may be the same as (or similar to) the deposition and coating conditions for the hole injection layer.

In some embodiments, the electron transport layer may include at least one compound selected from a compound represented by Formula 601 and a compound represented by Formula 602 illustrated below:



In Formula 601,

Ar₆₀₁ may be selected from the group consisting of: a naphthalene, a heptalene, a fluorene, a spiro-fluorene, a benzofluorene, a dibenzofluorene, a phenalene, a phenanthrene, an anthracene, a fluoranthene, a triphenylene, a pyrene, a chrysene, a naphthacene, a picene, a perylene, a pentaphene, and an indenoanthracene, and

a naphthalene group, a heptalene group, a fluorene group, a spiro-fluorene group, a benzofluorene group, a dibenzofluorene group, a phenalene group, a phenanthrene group, an anthracene group, a fluoranthene group, a triphenylene group, a pyrene group, a chrysene group, a naphthacene group, a picene group, a perylene group, a pentaphene group, and an indenoanthracene group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, and —Si(Q₃₀₁)(Q₃₀₂)(Q₃₀₃) (wherein Q₃₀₁ to Q₃₀₃ may each independently be selected from hydrogen, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ aryl group, and a C₁-C₆₀ heteroaryl group); description of L₆₀₁ may be the same as the description provided in connection with L₂₀₁;

E₆₀₁ may be selected from the group consisting of:

a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, an indolyl group, an indazolyl group, a purinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a phthalazinyl group, a naphthyridinyl group, a quinoxalinyl group, a quinazolinyl group, a cinnolinyl group, a carbazolyl group, a phenanthridinyl group, an acridinyl group, a phenanthroline group, a phenazinyl group, benzoimidazolyl, a benzofuranyl group, a benzothiophenyl group, isobenzothiazolyl, a benzoxazolyl group, an isobenzoxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group, and

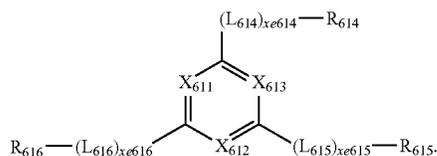
a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, an indolyl group, an indazolyl group, a purinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a phthalazinyl group, a naphthyridinyl group, a quinoxalinyl group, a quinazolinyl group, a cinnolinyl group, a carbazolyl group, a phenanthridinyl group, an acridinyl group, a phenanthroline group, a phenazinyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof,

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a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an azulenyl group, a heptalenyl group, an indacenyl group, an acenaphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenalenyl group, a phenanthrenyl group, an anthracenyl group, a fluoranthenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a naphthacenyl group, a picenyl group, a perylenyl group, a pentaphenyl group, a hexacenyl group, a pentacenyl group, a rubicenyl group, a coronenyl group, an ovalenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, an indolyl group, an indazolyl group, a purinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a phthalazinyl group, a naphthyridinyl group, a quinoxalinyl group, a quinazolinyl group, a cinnolinyl group, a carbazolyl group, a phenanthridinyl group, an acridinyl group, a phenanthrolinyl group, a phenazinyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, an imidazopyrimidinyl group, and an imidazopyrimidinyl group;

xe1 may be selected from 0, 1, 2, and 3; and

xe2 may be selected from 1, 2, 3, and 4.



Formula 602

In Formula 602,

X₆₁₁ may be N or C-(L₆₁₁)_{xe611}-R₆₁₁, X₆₁₂ may be N or C-(L₆₁₂)_{xe612}-R₆₁₂, X₆₁₃ may be N or C-(L₆₁₃)_{xe613}-R₆₁₃, and at least one selected from X₆₁₁ to X₆₁₃ may be N;

descriptions of L₆₁₁ to L₆₁₆ may each independently be the same as the description provided in connection with L₁;

R₆₁₁ to R₆₁₆ may each independently be selected from the group consisting of:

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group, and

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group, each substituted with at least one selected from deuterium, —F,

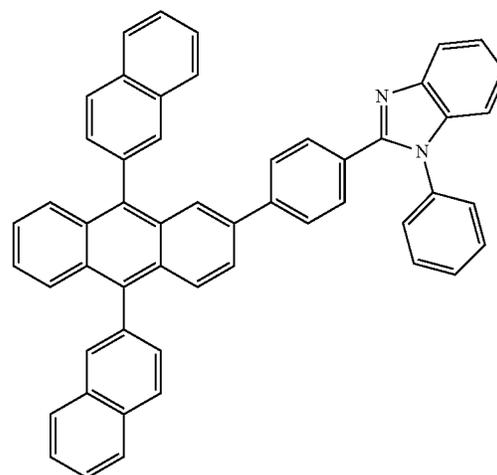
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—Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, an azulenyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group; and

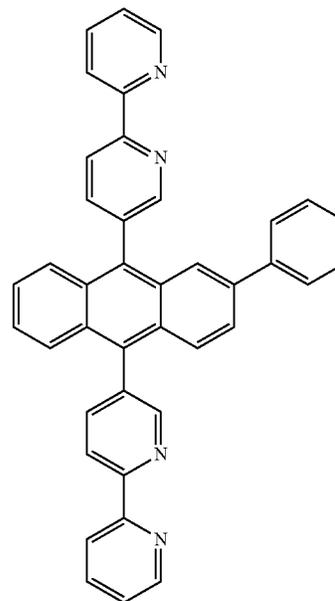
xe611 to xe616 may each independently be selected from 0, 1, 2, and 3.

The compound represented by Formula 601 and the compound represented by Formula 602 may each independently be selected from Compounds ET1 to ET15 illustrated below:

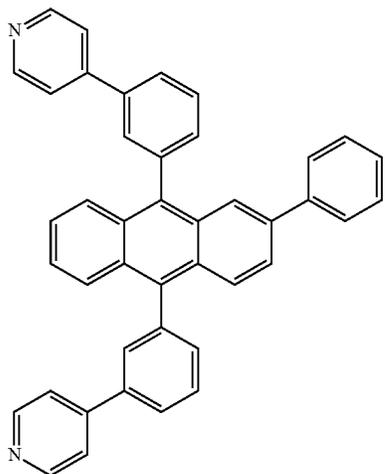
ET1



ET2



127
-continued



ET3

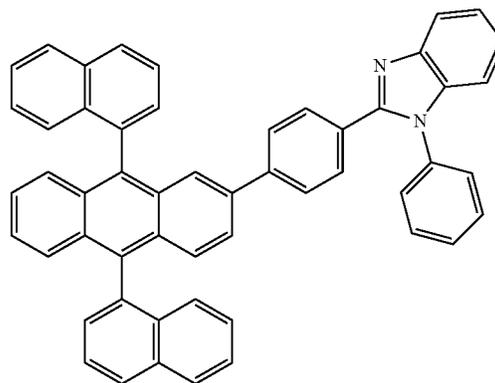
5

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15

20

128
-continued



ET6

25

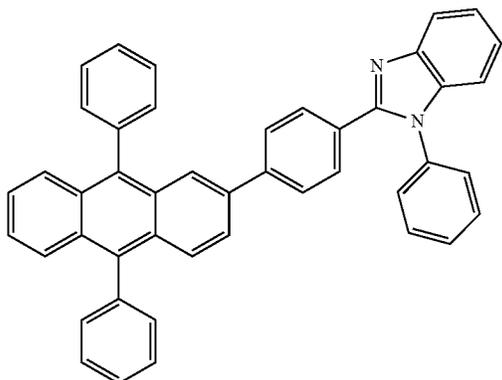
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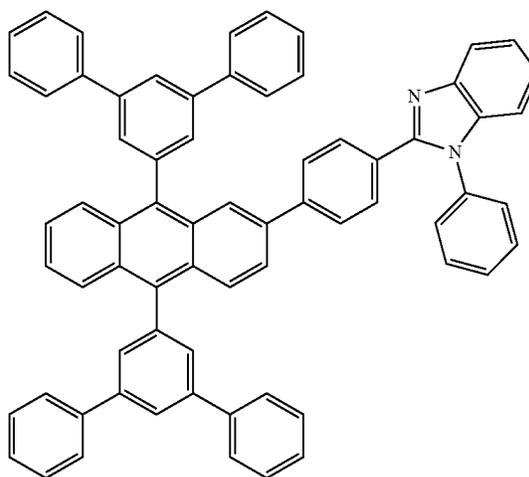
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ET4



ET7



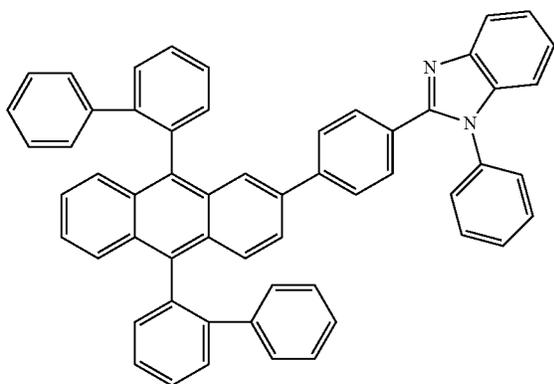
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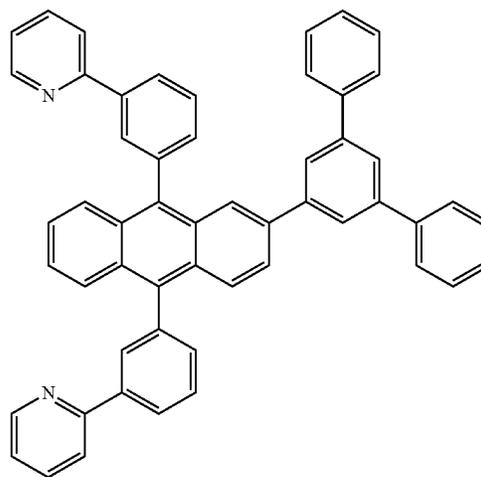
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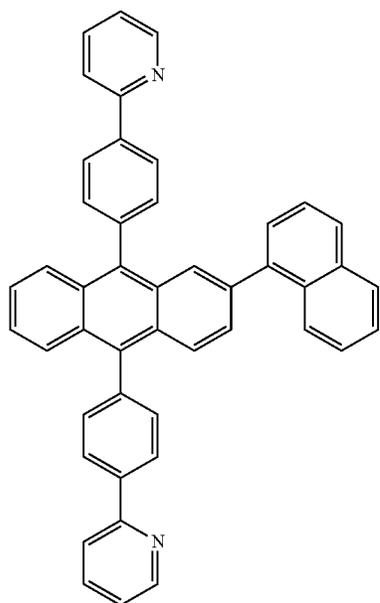
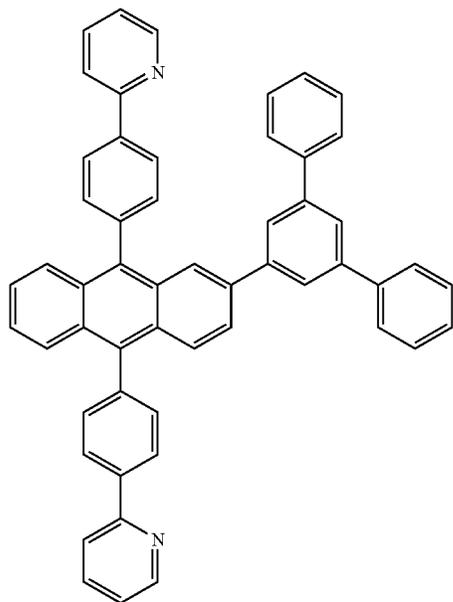
ET5



ET8



129
-continued



130
-continued

ET9

ET11

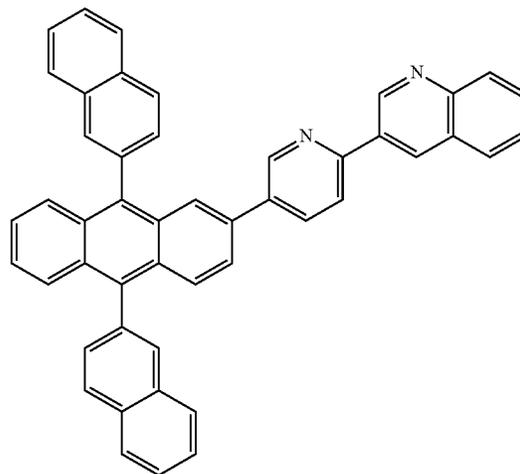
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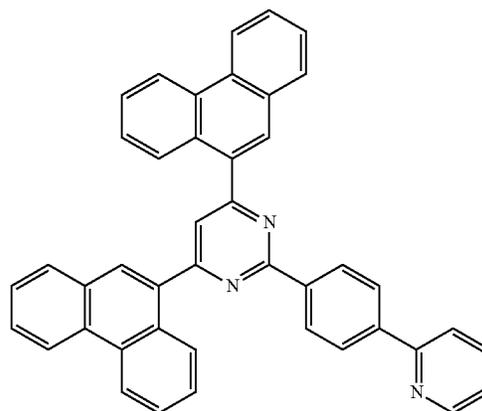
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ET10

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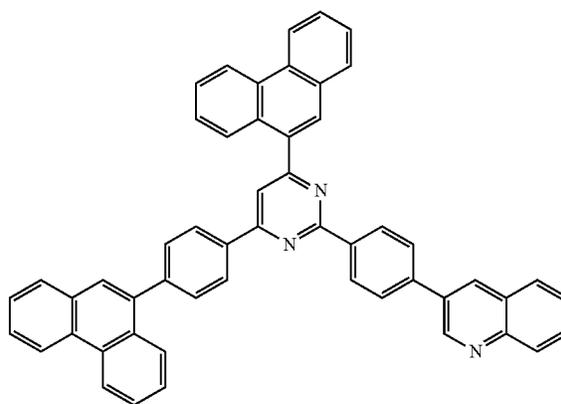
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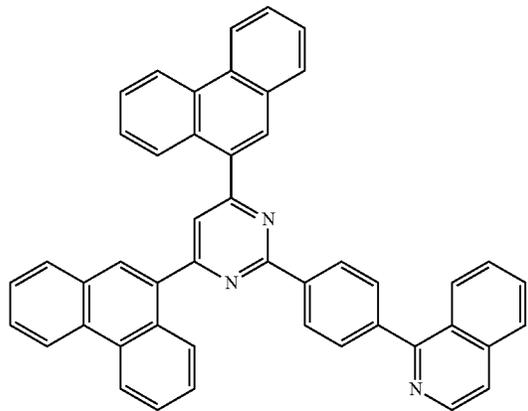
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ET12

ET13

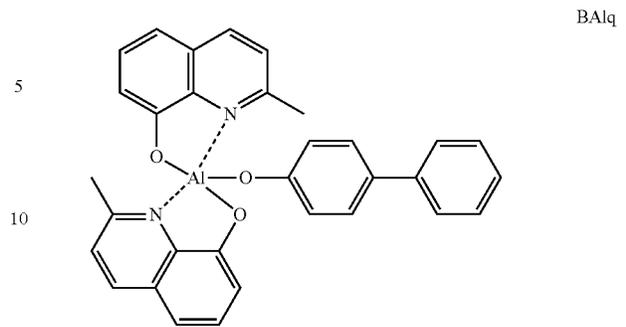


131
-continued



ET14

132
-continued



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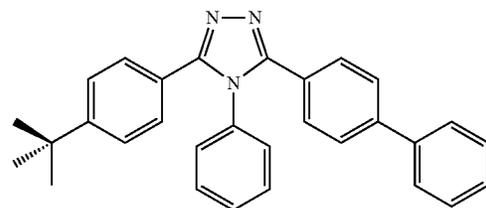
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BAlq

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TAZ

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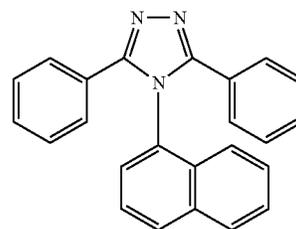


ET15

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NTAZ

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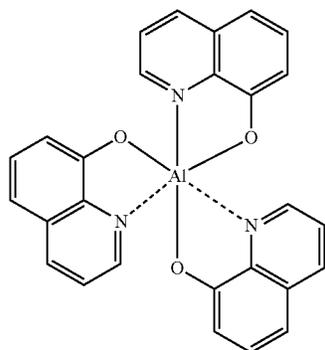
In some embodiments, the electron transport layer may further include at least one selected from BCP, Bphen, Alq₃, Balq, TAZ, and NTAZ.

A thickness of the electron transport layer may be in a range of about 100 Å to about 1,000 Å, for example, about 150 Å to about 500 Å. When the thickness of the electron transport layer is within any of the ranges described above, the electron transport layer may have satisfactory (or suitable) electron transport characteristics without a substantial increase in driving voltage.

Alq₃

The electron transport layer may further include, in addition to the materials described above, a metal-containing material.

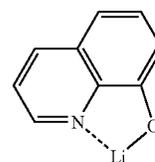
The metal-containing material may include a Li complex. The Li complex may include, for example, Compound ET-D1 (lithium quinolate, LiQ) and/or Compound ET-D2.



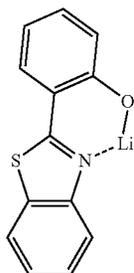
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ET-D1

65



-continued



The electron transport region may include an electron injection layer that may function to facilitate injection of electrons from the second electrode **190**.

The electron injection layer may be formed on the electron transport layer by using one or more suitable methods such as vacuum deposition, spin coating, casting, a LB method, ink-jet printing, laser-printing, and/or laser-induced thermal imaging. When an electron injection layer is formed by vacuum deposition and/or spin coating, deposition and coating conditions for the electron injection layer may be the same as (or similar to) those for the hole injection layer.

The electron injection layer may include at least one selected from LiF, NaCl, CsF, Li₂O, BaO, and LiQ.

A thickness of the electron injection layer may be in a range of about 1 Å to about 100 Å, for example, about 3 Å to about 90 Å. When the thickness of the electron injection layer is within any of the ranges described above, the electron injection layer may have satisfactory (or suitable) electron injection characteristics without a substantial increase in driving voltage.

The second electrode **190** may be disposed (e.g., positioned) on the organic layer **150** having the structure according to embodiments of the present disclosure. The second electrode **190** may be a cathode, which is an electron injection electrode, and in this regard, a material for forming the second electrode **190** may be selected from a metal, an alloy, an electrically conductive compound, and a mixture thereof, which have a relatively low work function. Non-limiting examples of the material for forming the second electrode **190** include lithium (Li), magnesium (Mg), aluminum (Al), aluminum-lithium (Al—Li), calcium (Ca), magnesium-indium (Mg—In), and magnesium-silver (Mg—Ag). In some embodiments, the material for forming the second electrode **190** may be ITO and/or IZO. The second electrode **190** may be a semi-transmissive electrode or a transmissive electrode.

Hereinbefore, the organic light-emitting device has been described with reference to the drawing, but embodiments of the present disclosure are not limited thereto.

The term “C₁-C₆₀ alkyl group,” as used herein, may refer to a linear or branched aliphatic hydrocarbon monovalent group having 1 to 60 carbon atoms, and non-limiting examples thereof include a methyl group, an ethyl group, a propyl group, an isobutyl group, a sec-butyl group, a tert-butyl group, a pentyl group, an iso-amyl group, and a hexyl group. The term “C₁-C₆₀ alkylene group,” as used herein, may refer to a divalent group having the same structure as the C₁-C₆₀ alkyl group.

The term “C₁-C₆₀ alkoxy group,” as used herein, may refer to a monovalent group represented by —OA₁₀₁ (wherein A₁₀₁ is the C₁-C₆₀ alkyl group), and non-limiting examples thereof include a methoxy group, an ethoxy group, and an isopropoxy group.

ET-D2

The term “C₂-C₆₀ alkenyl group,” as used herein, may refer to a hydrocarbon group having at least one carbon double bond at one or more positions along the hydrocarbon chain of the C₂-C₆₀ alkyl group (e.g., in the middle or at either terminal end of the C₂-C₆₀ alkyl group), and non-limiting examples thereof include an ethenyl group, a propenyl group, and a butenyl group. The term “C₂-C₆₀ alkenylene group,” as used herein, may refer to a divalent group having the same structure as the C₂-C₆₀ alkenyl group.

The term “C₂-C₆₀ alkynyl group,” as used herein, may refer to a hydrocarbon group having at least one carbon triple bond at one or more positions along the hydrocarbon chain of the C₂-C₆₀ alkyl group (e.g., in the middle or at either terminal end of the C₂-C₆₀ alkyl group), and non-limiting examples thereof include an ethynyl group and a propynyl group. The term “C₂-C₆₀ alkynylene group,” as used herein, may refer to a divalent group having the same structure as the C₂-C₆₀ alkynyl group.

The term “C₃-C₁₀ cycloalkyl group,” as used herein, may refer to a monovalent hydrocarbon monocyclic group having 3 to 10 carbon atoms, and non-limiting examples thereof include a cyclopropyl group, a cyclobutyl group, a cyclopentyl group, a cyclohexyl group, and a cycloheptyl group. The term “C₃-C₁₀ cycloalkylene group,” as used herein, may refer to a divalent group having the same structure as the C₃-C₁₀ cycloalkyl group.

The term “C₁-C₁₀ heterocycloalkyl group,” as used herein, may refer to a monovalent monocyclic group having at least one heteroatom selected from N, O, Si, P, and S as a ring-forming atom, and 1 to 10 carbon atoms, and non-limiting examples thereof include a tetrahydrofuranlyl group and a tetrahydrothiophenyl group. The term “C₁-C₁₀ heterocycloalkylene group,” as used herein, may refer to a divalent group having the same structure as the C₁-C₁₀ heterocycloalkyl group.

The term “C₃-C₁₀ cycloalkenyl group,” as used herein, may refer to a monovalent monocyclic group that has 3 to 10 carbon atoms and at least one double bond in the ring thereof and does not have aromaticity, and non-limiting examples thereof include a cyclopentenyl group, a cyclohexenyl group, and a cycloheptenyl group. The term “C₃-C₁₀ cycloalkenylene group,” as used herein, may refer to a divalent group having the same structure as the C₃-C₁₀ cycloalkenyl group.

The term “C₁-C₁₀ heterocycloalkenyl group,” as used herein, may refer to a monovalent monocyclic group that has at least one heteroatom selected from N, O, Si, P, and S as a ring-forming atom, 1 to 10 carbon atoms, and at least one double bond in its ring. Non-limiting examples of the C₁-C₁₀ heterocycloalkenyl group include a 2,3-hydrofuranlyl group and a 2,3-hydrothiophenyl group. The term “C₁-C₁₀ heterocycloalkenylene group,” as used herein, may refer to a divalent group having the same structure as the C₁-C₁₀ heterocycloalkenyl group.

The term “C₆-C₆₀ aryl group,” as used herein, may refer to a monovalent group having a carbocyclic aromatic system having 6 to 60 carbon atoms, and the term “C₆-C₆₀ arylene group,” as used herein, may refer to a divalent group having a carbocyclic aromatic system having 6 to 60 carbon atoms. Non-limiting examples of the C₆-C₆₀ aryl group include a phenyl group, a naphthyl group, an anthracenyl group, a phenanthrenyl group, a pyrenyl group, and a chrysenyl group. When the C₆-C₆₀ aryl group and the C₆-C₆₀ arylene group each independently include two or more rings, the respective rings may be fused to each other.

The term “C₁-C₆₀ heteroaryl group,” as used herein, may refer to a monovalent group having a carbocyclic aromatic

system that has at least one heteroatom selected from N, O, Si, P, and S as a ring-forming atom, and 1 to 60 carbon atoms. The term “C₁-C₆₀ heteroarylene group,” as used herein, may refer to a divalent group having a carbocyclic aromatic system that has at least one heteroatom selected from N, O, P, and S as a ring-forming atom, and 1 to 60 carbon atoms. Non-limiting examples of the C₁-C₆₀ heteroaryl group include a pyridinyl group, a pyrimidinyl group, a pyrazinyl group, a pyridazinyl group, a triazinyl group, a quinolinyl group, and an isoquinolinyl group. When the C₁-C₆₀ heteroaryl group and the C₁-C₆₀ heteroarylene group each independently include two or more rings, the respective rings may be fused to each other.

The term “C₆-C₆₀ aryloxy group,” as used herein, may refer to a monovalent group represented by —OA₁₀₂ (wherein A₁₀₂ is the C₆-C₆₀ aryl group), and the term “C₆-C₆₀ arylthio group,” as used herein, may refer to a monovalent group represented by —SA₁₀₃ (wherein A₁₀₃ is the C₆-C₆₀ aryl group).

The term “monovalent non-aromatic condensed polycyclic group,” as used herein, may refer to a monovalent group that has two or more rings condensed (e.g., fused) to each other, only carbon atoms as ring forming atoms (e.g., 8 to 60 carbon atoms), and non-aromaticity in the entire molecular structure (e.g., not having overall aromaticity). A non-limiting example of the monovalent non-aromatic condensed polycyclic group is a fluorenyl group. The term “divalent non-aromatic condensed polycyclic group,” as used herein, may refer to a divalent group having the same structure as the monovalent non-aromatic condensed polycyclic group.

The term “monovalent non-aromatic condensed heteropolycyclic group,” as used herein, may refer to a monovalent group that has two or more rings condensed (e.g., fused) to each other, has at least one heteroatom selected from N, O, Si, P, and S, other than carbon atoms (e.g., 2 to 60 carbon atoms), as ring forming atoms, and has non-aromaticity in the entire molecular structure (e.g., does not have overall aromaticity). Non-limiting example of the monovalent non-aromatic condensed heteropolycyclic group is a carbazolyl group. The term “divalent non-aromatic condensed heteropolycyclic group,” as used herein, may refer to a divalent group having the same structure as the monovalent non-aromatic condensed heteropolycyclic group.

At least one substituent of the substituted fluorenyl group, substituted spiro-bifluorenyl group, substituted carbazolyl group, substituted dibenzofuranyl group, substituted dibenzothiofenyl group, substituted dibenzosilolyl group, substituted C₃-C₁₀ cycloalkylene group, substituted C₁-C₁₀ heterocycloalkylene group, substituted C₃-C₁₀ cycloalkenylene group, substituted C₁-C₁₀ heterocycloalkenylene group, substituted C₆-C₆₀ arylene group, substituted C₁-C₆₀ heteroarylene group, substituted divalent non-aromatic condensed polycyclic group, substituted divalent non-aromatic condensed heteropolycyclic group, substituted C₁-C₆₀ alkyl group, substituted C₂-C₆₀ alkenyl group, substituted C₂-C₆₀ alkynyl group, substituted C₁-C₆₀ alkoxy group, substituted C₃-C₁₀ cycloalkyl group, substituted C₁-C₁₀ heterocycloalkyl group, substituted C₃-C₁₀ cycloalkenyl group, substituted C₁-C₁₀ heterocycloalkenyl group, substituted C₆-C₆₀ aryl group, substituted C₆-C₆₀ aryloxy group, substituted C₆-C₆₀ arylthio group, substituted C₁-C₆₀ heteroaryl group, substituted monovalent non-aromatic condensed polycyclic group, and substituted monovalent non-aromatic condensed heteropolycyclic group may be selected from the group consisting of:

deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group,

a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group (e.g., aryloxy), C₆-C₆₀ arylthio group (e.g., arylthio), C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, and —Si(Q₁₁)(Q₁₂)(Q₁₃),

a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, a biphenyl group, and a terphenyl group,

a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, a biphenyl group, and a terphenyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, a biphenyl group, and a terphenyl group, and —Si(Q₂₁)(Q₂₂)(Q₂₃), and

—Si(Q₃₁)(Q₃₂)(Q₃₃),
wherein Q₁₁ to Q₁₃, Q₂₁ to Q₂₃, and Q₃₁ to Q₃₃ may each independently be selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, a biphenyl group, and a terphenyl group.

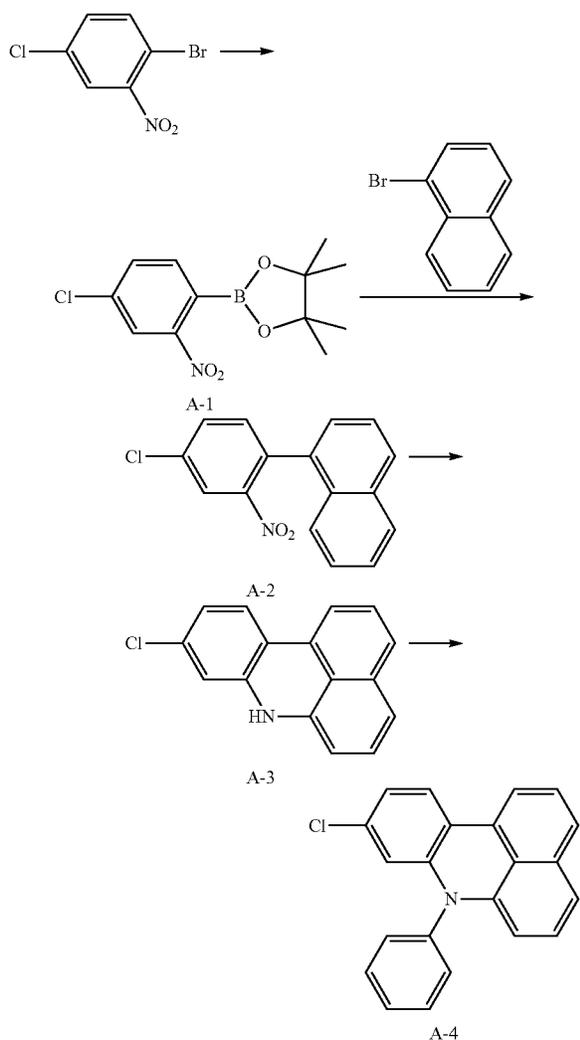
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The term "Ph," used herein, may refer to a phenyl group, the term "Me," used herein, may refer to a methyl group, the term "Et," used herein, may refer to an ethyl group, the term "ter-Bu," or "But" used herein, may refer to a tert-butyl group, and "D" may refer to deuterium.

Hereinafter, an organic light-emitting device according to one or more embodiments of the present disclosure will be described in further detail with reference to Synthesis Examples and Examples. The expression "B was used instead of A" used in describing Synthesis Examples may refer to a molar equivalent of A being identical to a molar equivalent of B.

EXAMPLES

Synthesis Example 1: Synthesis of Compound 10



Synthesis of Intermediate A-1

5.96 g (23.6 millimoles (mmol)) of 1-bromo-4-chloro-2-nitrobenzene was dissolved in 200 milliliters (mL) of THF, and 10 mL (25.0 mmol, 2.5M in Hexane) of n-BuLi was slowly added dropwise thereto at a temperature of -78°C . After 1 hour of stirring the resulting solution at the same temperature, 9.3 mL (50 mmol) of 2-isopropoxy-4,4,5,5-

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tetramethyl-1,3,2-dioxaborolane was slowly added dropwise thereto, and then, the obtained reaction solution was stirred at a temperature of -78°C . for 1 hour. The resulting solution was additionally stirred for 24 hours at room temperature, and once the reaction was completed, 50 mL of a 10% HCl aqueous solution and 50 mL of H_2O were added thereto, followed by three times of extraction using 80 mL of diethylether. An organic layer collected from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 6.36 grams (g) (yield: 90%) of Intermediate A-1. The prepared compound was identified by liquid chromatography-mass spectroscopy (LC-MS).

$(\text{C}_{12}\text{H}_{15}\text{BClNO}_4: \text{M}+ 283.08)$

Synthesis of Intermediate A-2

6.23 g (22.0 mmol) of Intermediate A-1, 4.56 g (22.0 mmol) of 1-bromonaphthalene, 1.27 g (1.1 mmol) of tetrakis (triphenylphosphine)palladium ($\text{Pd}(\text{PPh}_3)_4$), and 4.5 g (33 mmol) of K_2CO_3 were dissolved in 200 mL of a THF/ H_2O (2/1 volume ratio) mixed solution, and then, the resulting solution was stirred at a temperature of 70°C . for 5 hours. After the obtained reaction solution was cooled to room temperature, 60 mL of water was added thereto, followed by three times of extraction using 60 mL of ethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and then, a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 4.68 g (yield: 75%) of Intermediate A-2. The prepared compound was identified by LC-MS.

$(\text{C}_{16}\text{H}_{10}\text{ClNO}_2: \text{M}+ 283.04)$

Synthesis of Intermediate A-3

2.83 g (10.00 mmol) of Intermediate A-2 and 8.3 g (50 mmol) of triethylphosphite were dissolved in 250 mL of cumene, and then, the resulting mixed solution was stirred under reflux in a nitrogen atmosphere for 24 hours. After the reaction was completed, an organic layer obtained by concentration under reduced pressure in high vacuum was separation-purified by silica gel column chromatography, thereby completing the preparation of 1.76 g (yield: 70%) of Intermediate A-3. The prepared compound was identified by LC-MS.

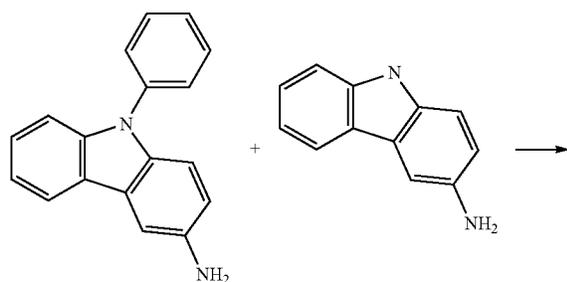
$(\text{C}_{16}\text{H}_{10}\text{ClN}: \text{M}+ 251.71)$

Synthesis of Intermediate A-4

1.26 g (5.00 mmol) of Intermediate A-3, 0.95 g (6 mmol) of bromobenzene, 0.003 g (0.1 mmol) of $\text{Pd}(\text{OAc})_2$, 0.1 g (0.5 mmol) of $\text{P}(\text{tBu})_3$, and 0.84 g (7.5 mmol) of KOtBu were dissolved in 100 mL of toluene, and then, the resulting mixed solution was stirred under reflux in a nitrogen atmosphere for about 24 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon three times using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 1.31 g (yield: 80%) of Intermediate A-4. The prepared compound was identified by LC-MS.

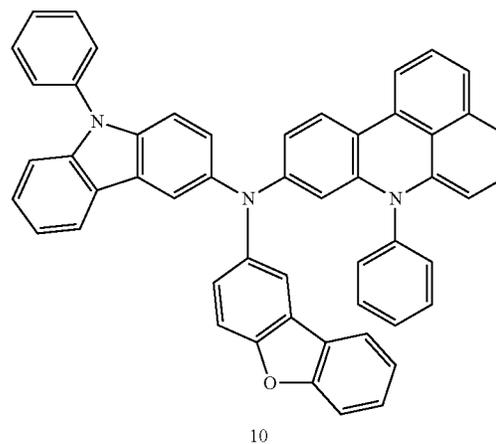
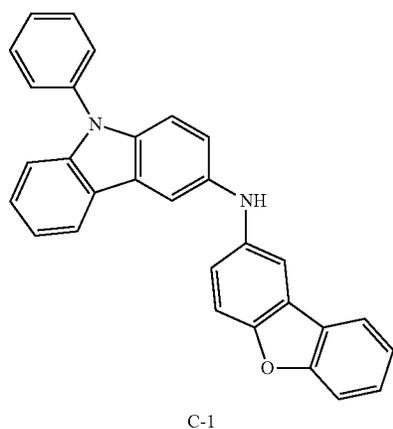
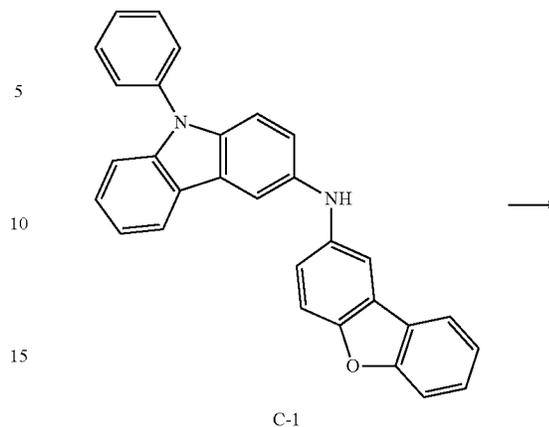
$(\text{C}_{22}\text{H}_{14}\text{ClN}: \text{M}+ 327.81)$

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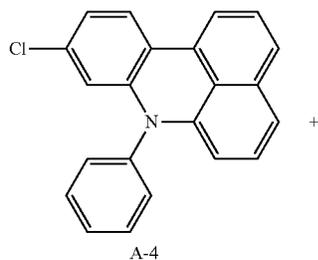
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Synthesis of Intermediate C-1

7.75 g (30 mmol) of 9-phenyl-9H-carbazol-3-amine, 4.94 g (20.0 mmol) of 2-bromodibenzo[b,d]furan, 0.37 g (0.4 mmol) of $\text{Pd}_2(\text{dba})_3$, 0.08 g (0.4 mmol) of $\text{P}(\text{tBu})_3$, and 5.76 g (60 mmol) of KOtBu were dissolved in 200 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 6.79 g (yield: 80%) of Intermediate C-1. The prepared compound was identified by LC-MS.

($\text{C}_{30}\text{H}_{20}\text{N}_2\text{O}$: M^+ 424.50)



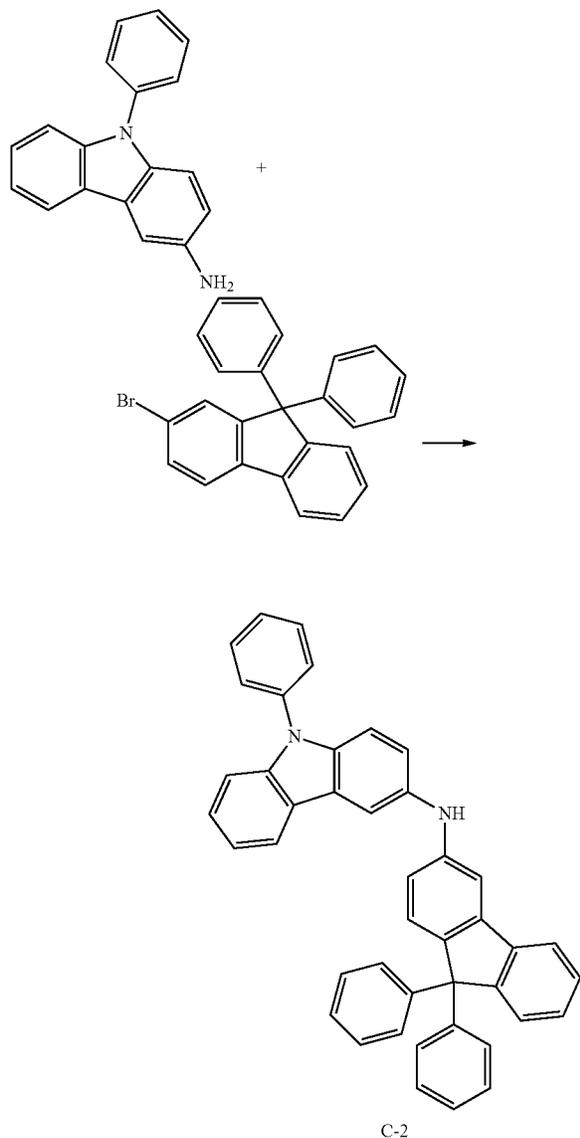
Synthesis of Compound 10

1.64 g (5.00 mmol) of Intermediate A-4, 2.12 g (5 mmol) of Intermediate C-1, 0.093 g (0.1 mmol) of $\text{Pd}_2(\text{dba})_3$, 0.1 g (0.5 mmol) of $\text{P}(\text{tBu})_3$, and 0.84 g (7.5 mmol) of KOtBu were dissolved in 100 mL of toluene, and then, the resulting mixed solution was stirred under reflux in a nitrogen atmosphere for 24 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon three times using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 2.97 g (yield: 83%) of Compound 10. The prepared compound was identified by LC-MS and nuclear magnetic resonance (NMR).

($\text{C}_{52}\text{H}_{33}\text{N}_3\text{O}$: M^+ 715.86)

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Synthesis Example 2: Synthesis of Compound 14

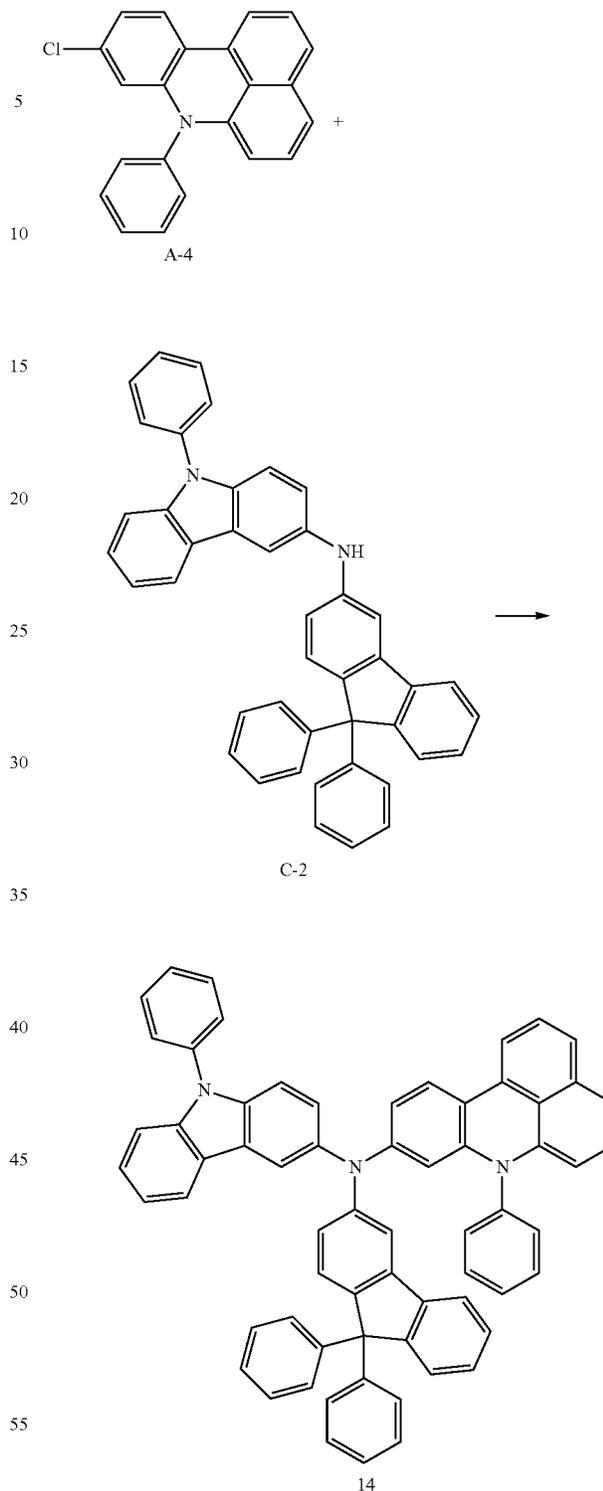


Synthesis of Intermediate C-2

7.75 g (30 mmol) of 9-phenyl-9H-carbazol-3-amine, 7.95 g (20.0 mmol) of 2-bromo-9,9-diphenyl-9H-fluorene, 0.37 g (0.4 mmol) of $\text{Pd}_2(\text{dba})_3$, 0.08 g (0.4 mmol) of $\text{P}(\text{tBu})_3$, and 5.76 g (60 mmol) of KOtBu were dissolved in 200 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 9.31 g (yield: 81%) of Intermediate C-2. The prepared compound was identified by LC-MS.

($\text{C}_{43}\text{H}_{30}\text{N}_2$: M^+ 574.73)

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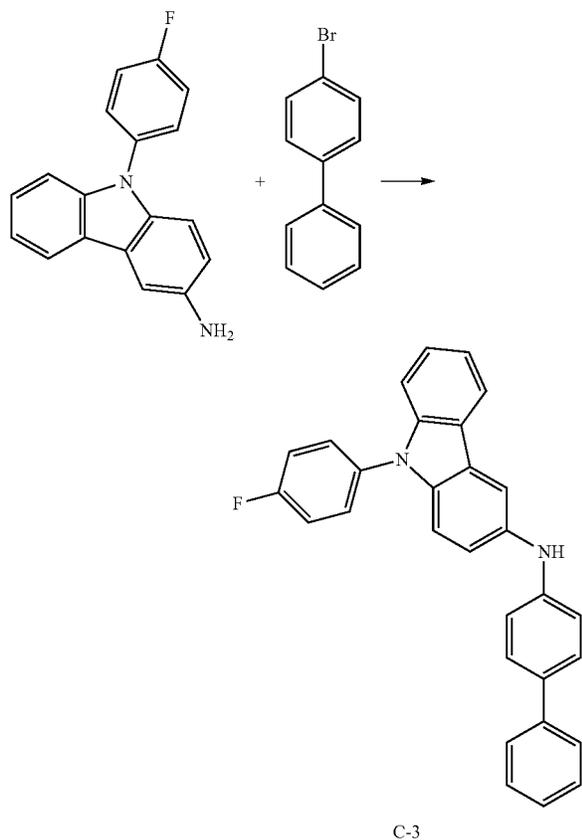
Synthesis of Compound 14

3.55 g (yield: 82%) of Compound 14 was obtained in the same (or substantially the same) manner as in Synthesis of Compound 10, except that 2.87 g (5 mmol) of Intermediate C-2 was used instead of Intermediate C-1. The prepared compound was identified by LC-MS and NMR.

($\text{C}_{65}\text{H}_{43}\text{N}_3$: M^+ 866.08)

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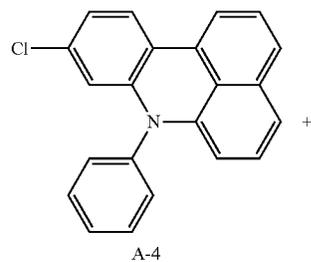
Synthesis Example 3: Synthesis of Compound 18



Synthesis of Intermediate C-3

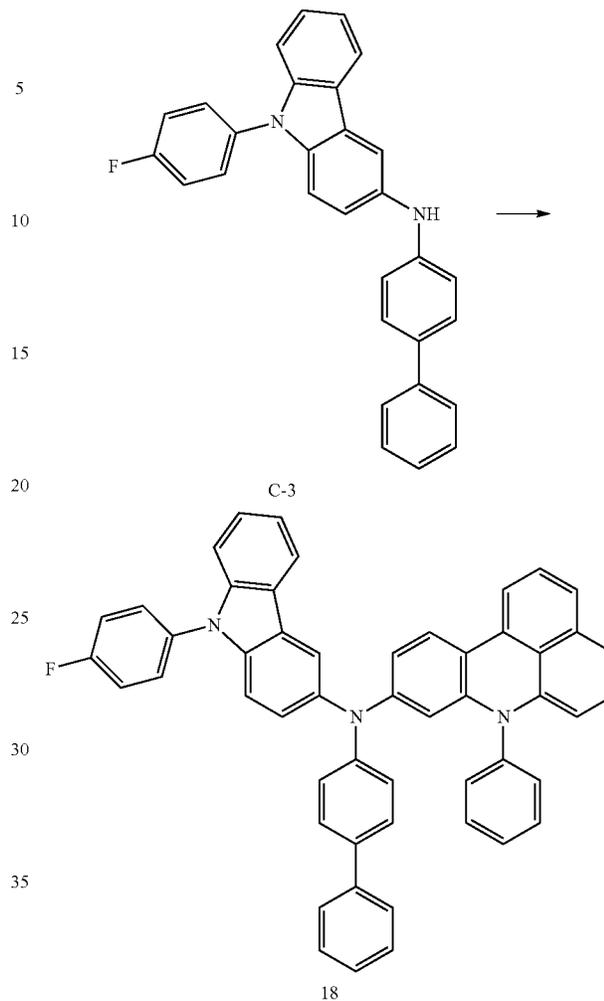
8.29 g (30 mmol) of 9-(4-fluorophenyl)-9H-carbazol-3-amine, 4.66 g (20.0 mmol) of 4-bromo-1,1'-biphenyl, 0.37 g (0.4 mmol) of $\text{Pd}_2(\text{dba})_3$, 0.08 g (0.4 mmol) of $\text{P}(\text{tBu})_3$, and 5.76 g (60 mmol) of KOtBu were dissolved in 200 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 6.86 g (yield: 80%) of Intermediate C-3. The prepared compound was identified by LC-MS.

($\text{C}_{30}\text{H}_{21}\text{FN}_2$: $\text{M}+$ 428.51)



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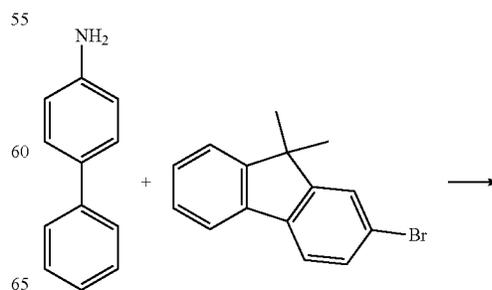


Synthesis of Compound 18

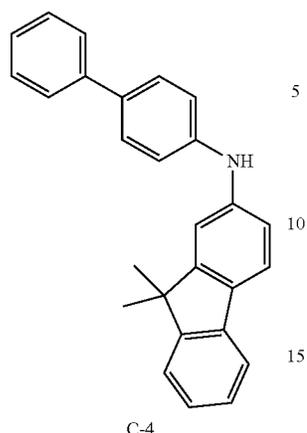
2.92 g (yield: 81%) of Compound 18 was obtained in the same (or substantially the same) manner as in Synthesis of Compound 10, except that 2.14 g (5 mmol) of Intermediate C-3 was used instead of Intermediate C-1. The prepared compound was identified by LC-MS and NMR.

($\text{C}_{52}\text{H}_{34}\text{FN}_3$: $\text{M}+$ 719.86)

Synthesis Example 4: Synthesis of Compound 22



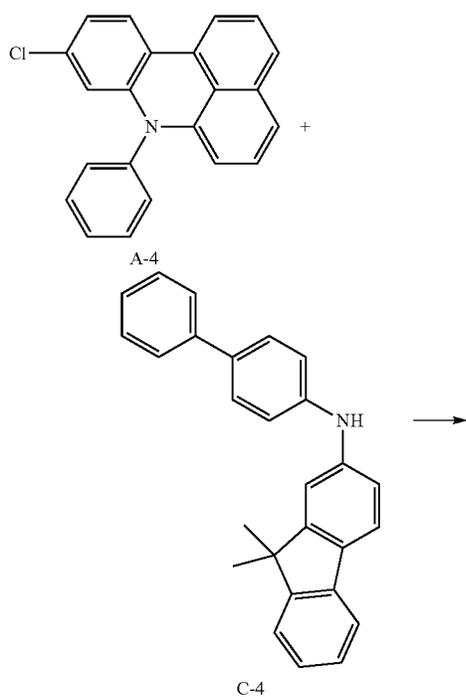
145
-continued



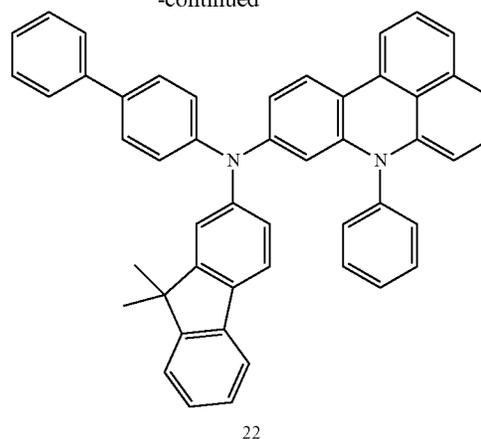
Synthesis of Intermediate C-4

5.08 g (30 mmol) of 4-aminobiphenyl, 5.46 g (20.0 mmol) of 2-bromo-9,9-dimethylfluorene, 0.37 g (0.4 mmol) of $\text{Pd}_2(\text{dba})_3$, 0.08 g (0.4 mmol) of $\text{P}(\text{tBu})_3$, and 5.76 g (60 mmol) of KOtBu were dissolved in 200 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 6.07 g (yield: 84%) of Intermediate C-4. The prepared compound was identified by LC-MS.

($\text{C}_{27}\text{H}_{23}\text{N}$: $M+$ 361.49)



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-continued

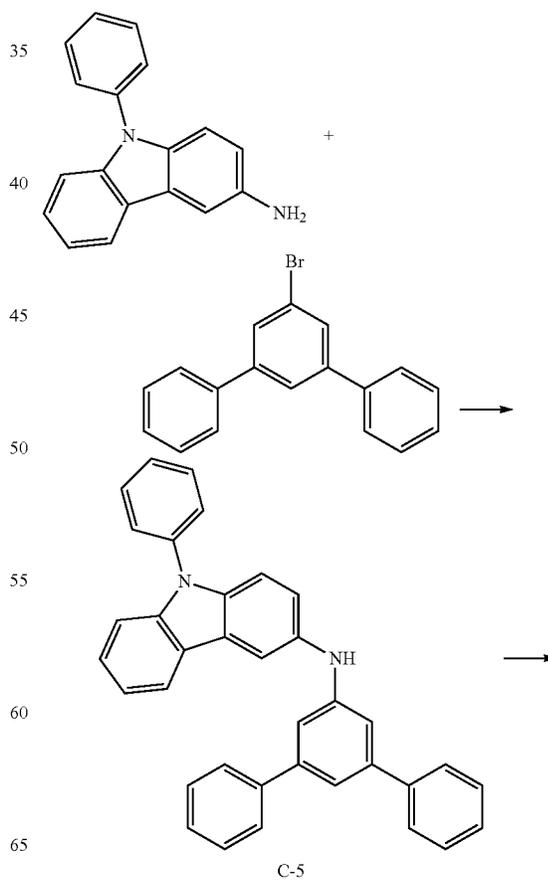


Synthesis of Compound 22

2.68 g (yield: 82%) of Compound 22 was obtained in the same (or substantially the same) manner as in Synthesis of Compound 10, except that 1.81 g (5 mmol) of Intermediate C-4 was used instead of Intermediate C-1. The prepared compound was identified by LC-MS and NMR.

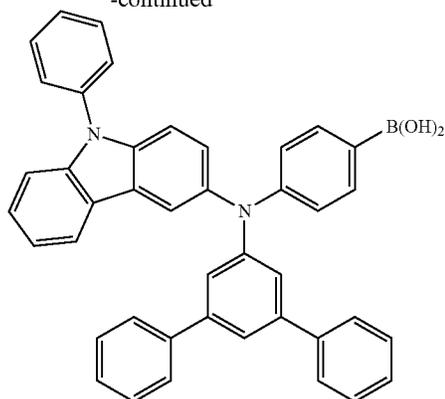
($\text{C}_{49}\text{H}_{36}\text{N}_2$: $M+$ 652.84)

Synthesis Example 5: Synthesis of Compound 29



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-continued



C-6

Synthesis of Intermediate C-5

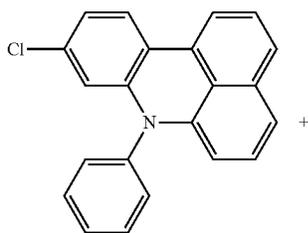
7.75 g (30 mmol) of 9-phenyl-9H-carbazol-3-amine, 6.18 g (20.0 mmol) of 5'-bromo-1,1':3',1''-terphenyl, 0.37 g (0.4 mmol) of $\text{Pd}_2(\text{dba})_3$, 0.08 g (0.4 mmol) of $\text{P}(\text{tBu})_3$, and 5.76 g (60 mmol) of KOtBu were dissolved in 200 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 7.69 g (yield: 79%) of Intermediate C-5. The prepared compound was identified by LC-MS.

($\text{C}_{36}\text{H}_{26}\text{N}_2$; M+ 486.62)

Synthesis of Intermediate C-6

9.73 g (20 mmol) of Intermediate C-5, 4.02 g (20.0 mmol) of (4-bromophenyl)boronic acid, 0.37 g (0.4 mmol) of $\text{Pd}_2(\text{dba})_3$, 0.08 g (0.4 mmol) of $\text{P}(\text{tBu})_3$, and 5.76 g (60 mmol) of KOtBu were dissolved in 200 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 7.69 g (yield: 79%) of Intermediate C-6. The prepared compound was identified by LC-MS.

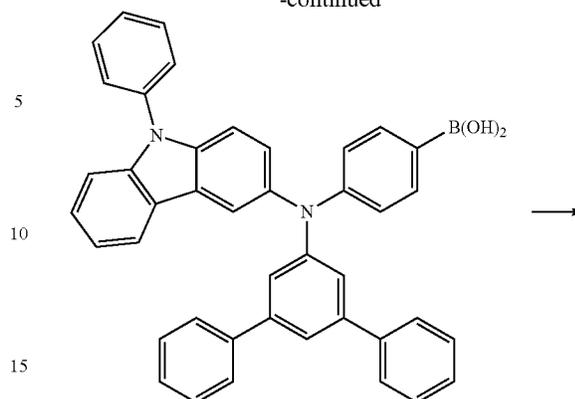
($\text{C}_{42}\text{H}_{31}\text{BN}_2\text{O}_2$; M+ 606.53)



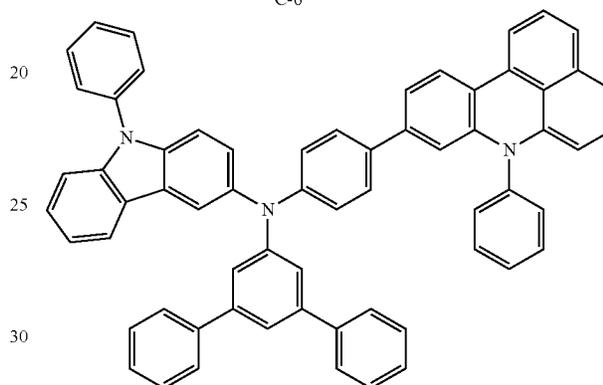
A-4

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C-6



29

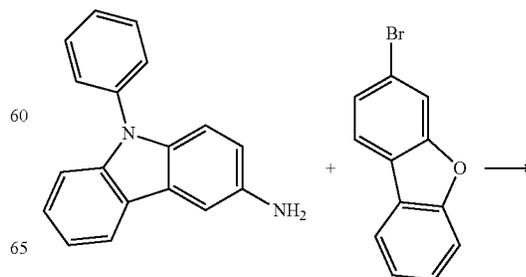
Synthesis of Compound 29

1.64 g (5.00 mmol) of Intermediate A-4, 2.43 g (5 mmol) of Intermediate C-6, 0.58 g (0.50 mmol) of $\text{Pd}(\text{PPh}_3)_4$, and 2.08 g (15 mmol) of K_2CO_3 were dissolved in 200 mL of a THF/ H_2O (2/1 volume ratio) mixed solution, and then, the resulting solution was stirred at a temperature of 70° C. for 5 hours. After the obtained reaction solution was cooled to room temperature, 60 mL of water was added thereto, followed by three times of extraction using 60 mL of ethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and then, a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 3.07 g (yield: 72%) of Compound 29. The prepared compound was identified by LC-MS and NMR.

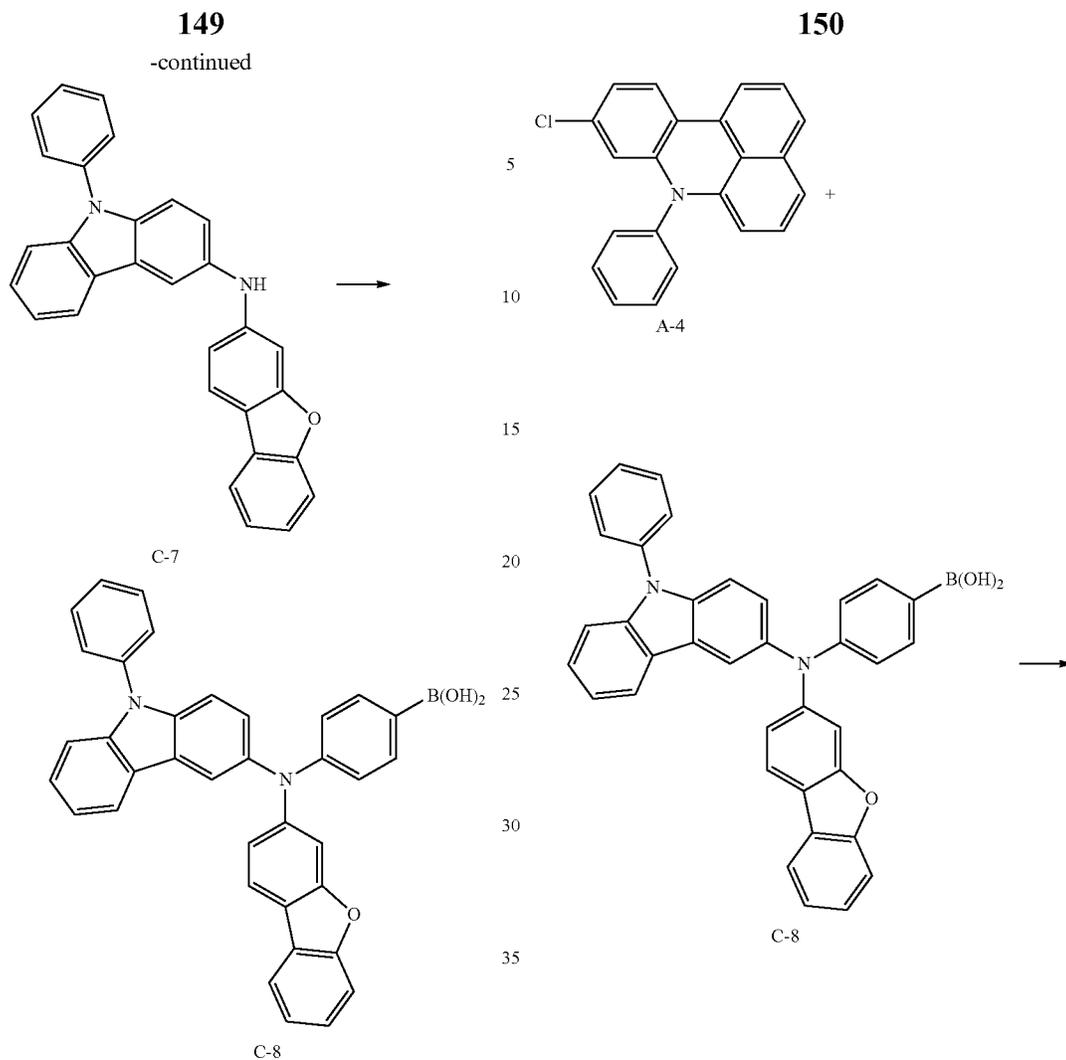
($\text{C}_{64}\text{H}_{43}\text{N}_3$; M+ 854.07)

Synthesis Example 6: Synthesis of Compound 36

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65



Synthesis of Intermediate C-7

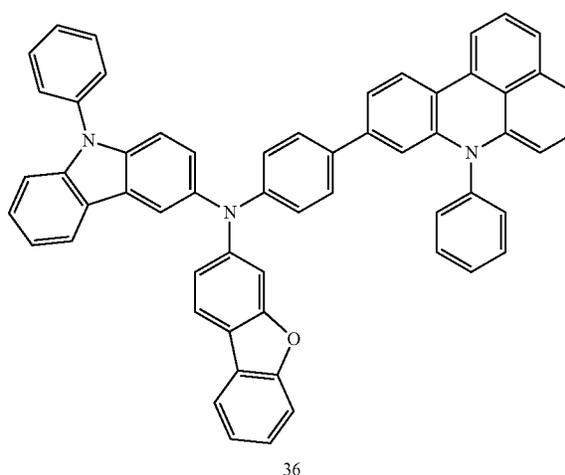
7.75 g (30 mmol) of 9-phenyl-9H-carbazol-3-amine, 4.94 g (20.0 mmol) of 3-bromodibenzo[b,d]furan, 0.37 g (0.4 mmol) of $\text{Pd}_2(\text{dba})_3$, 0.08 g (0.4 mmol) of $\text{P}(\text{tBu})_3$, and 5.76 g (60 mmol) of KOtBu were dissolved in 200 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 6.96 g (yield: 82%) of Intermediate C-7. The prepared compound was identified by LC-MS.

($\text{C}_{30}\text{H}_{20}\text{N}_2\text{O}$: $M+ 424.50$)

Synthesis of Intermediate C-8

8.49 g (yield: 78%) of Intermediate C-8 was obtained in the same (or substantially the same) manner as in Synthesis of Intermediate C-6, except that 12.12 g (20 mmol) of Intermediate C-7 was used instead of Intermediate C-5. The prepared compound was identified by LC-MS.

($\text{C}_{36}\text{H}_{25}\text{BN}_2\text{O}_3$: $M+ 544.42$)



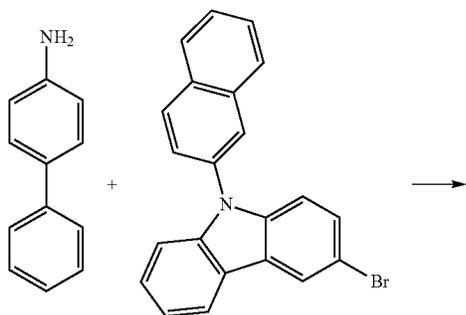
Synthesis of Compound 36

2.93 g (yield: 74%) of Compound 36 was obtained in the same (or substantially the same) manner as in Synthesis of Compound 29, except that 2.72 g (5 mmol) of Intermediate C-8 was used instead of Intermediate C-6. The prepared compound was identified by LC-MS and NMR.

($\text{C}_{58}\text{H}_{37}\text{N}_3\text{O}$: $M+ 791.95$)

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Synthesis Example 7: Synthesis of Compound 41



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Synthesis of Intermediate C-9

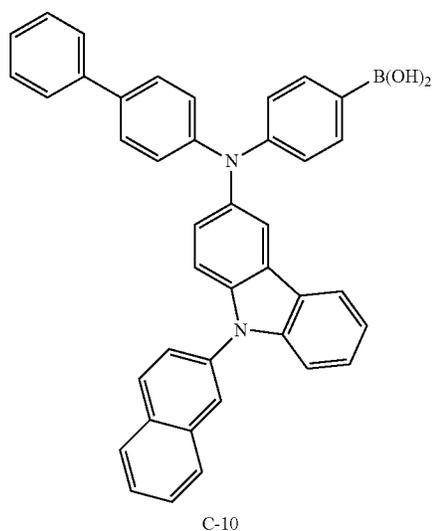
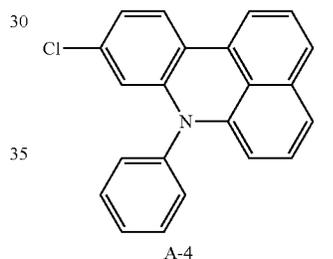
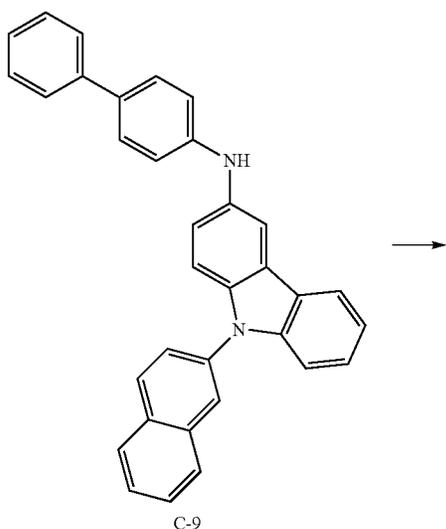
5.08 g (30 mmol) of 4-aminobiphenyl, 7.45 g (20.0 mmol) of 3-bromo-9-(naphthalen-2-yl)-9H-carbazole, 0.37 g (0.4 mmol) of $\text{Pd}_2(\text{dba})_3$, 0.08 g (0.4 mmol) of $\text{P}(\text{tBu})_3$, and 5.76 g (60 mmol) of KOtBu were dissolved in 200 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 7.36 g (yield: 80%) of Intermediate C-9. The prepared compound was identified by LC-MS.

($\text{C}_{34}\text{H}_{24}\text{N}_2$; M+ 460.58)

20 Synthesis of Intermediate C-10

9.06 g (yield: 78%) of Intermediate C-10 was obtained in the same (or substantially the same) manner as in Synthesis of Intermediate C-6, except that 9.21 g (20 mmol) of Intermediate C-9 was used instead of Intermediate C-5. The prepared compound was identified by LC-MS.

($\text{C}_{40}\text{H}_{29}\text{BN}_2\text{O}_2$; M+ 580.49)



40

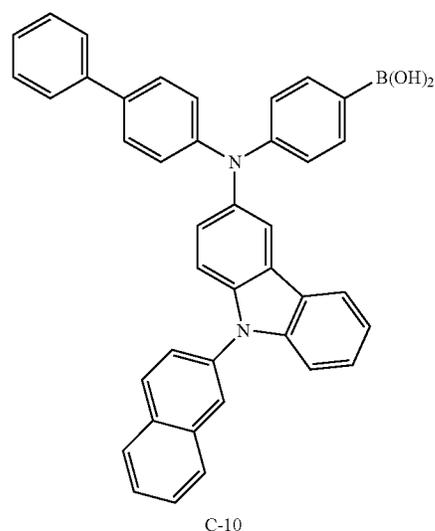
45

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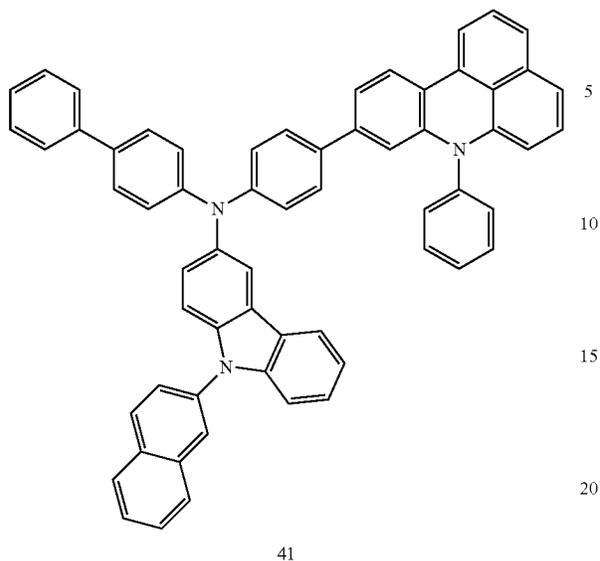
65



C-10

153

-continued

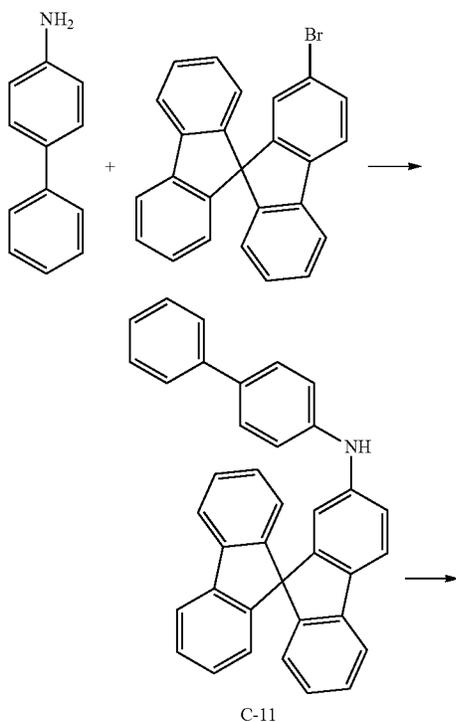


Synthesis of Compound 41

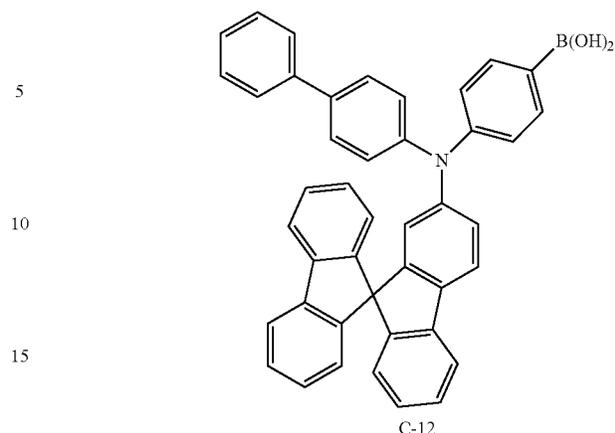
3.27 g (yield: 79%) of Compound 41 was obtained in the same (or substantially the same) manner as in Synthesis of Compound 29, except that 2.90 g (5 mmol) of Intermediate C-10 was used instead of Intermediate C-6. The prepared compound was identified by LC-MS and NMR.

(C₆₂H₄₁N₃: M+ 828.03)

Synthesis Example 8: Synthesis of Compound 48

**154**

-continued



Synthesis of Intermediate C-11

5.08 g (30 mmol) of 4-aminobiphenyl, 7.90 g (20.0 mmol) of 2-bromo-9,9'-spirobifluorene, 0.37 g (0.4 mmol) of Pd₂(dba)₃, 0.08 g (0.4 mmol) of P(tBu)₃, and 5.76 g (60 mmol) of KOtBu were dissolved in 200 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 7.25 g (yield: 75%) of Intermediate C-11. The prepared compound was identified by LC-MS.

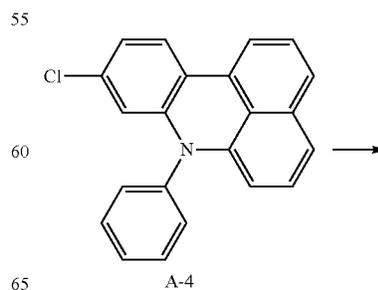
(C₃₇H₂₅N: M+ 483.61)

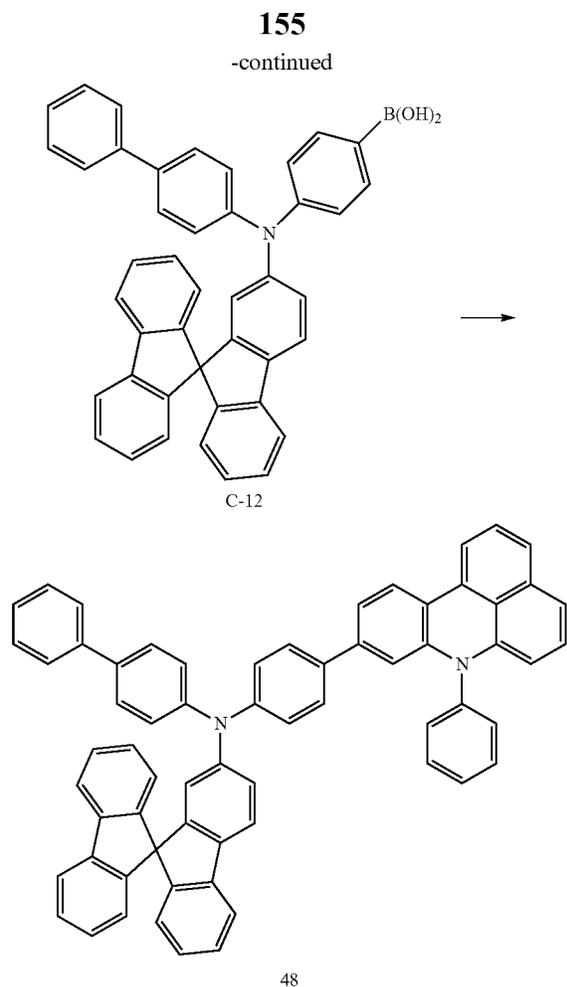
Synthesis of Intermediate C-12

8.93 g (yield: 74%) of Intermediate C-12 was obtained in the same (or substantially the same) manner as in Synthesis of Intermediate C-6, except that 9.67 g (20 mmol) of Intermediate C-11 was used instead of Intermediate C-5. The prepared compound was identified by LC-MS.

(C₄₃H₃₀BNO₂: M+ 603.53)

Synthesis of Compound 48

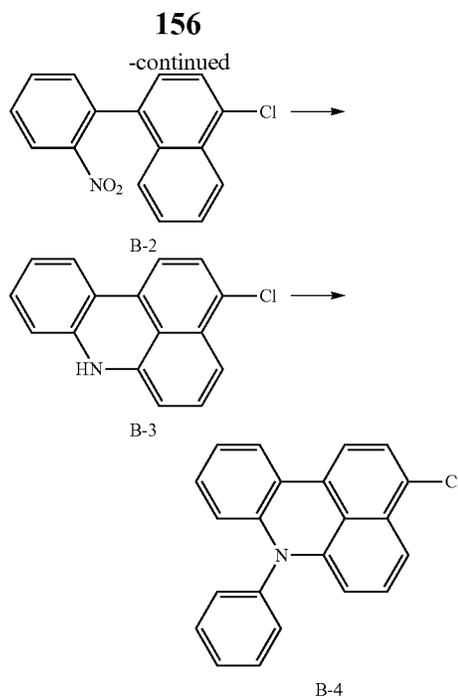
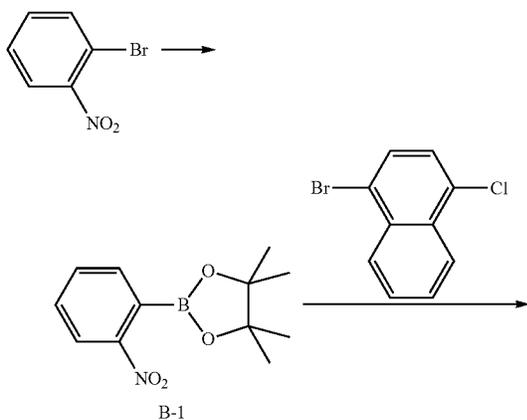




3.23 g (yield: 76%) of Compound 48 was obtained in the same (or substantially the same) manner as in Synthesis of Compound 29, except that 3.02 g (5 mmol) Intermediate C-12 was used instead of Intermediate C-6. The prepared compound was identified by LC-MS and NMR.

($C_{65}H_{42}N_2$: $M+$ 851.07)

Synthesis Example 9: Synthesis of Compound 54



Synthesis of Intermediate B-1

4.77 g (23.6 mmol) of 1-bromo-2-nitrobenzene was dissolved in 100 mL of THF, and then, 10 mL of n-BuLi (25.0 mmol, 2.5M in Hexane) was slowly added dropwise thereto at a temperature of $-78^{\circ}C$. After 1 hour of stirring the result at the same temperature, 9.3 mL (50 mmol) of 2-isopropoxy-4,4,5,5-tetramethyl-1,3,2-dioxaborolane was slowly added dropwise to the solution, and then, the obtained reaction solution was stirred at a temperature of $-78^{\circ}C$ for 1 hour. The resulting solution was additionally stirred for 24 hours at room temperature, and once the reaction was completed, 50 mL of a 10% HCl aqueous solution and 50 mL of H_2O were added thereto, followed by three times of extraction using 80 mL of diethylether. An organic layer collected from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 5.29 g (yield: 90%) of Intermediate B-1. The prepared compound was identified by LC-MS.

($C_{12}H_{16}BNO_4$: $M+$ 249.07)

Synthesis of Intermediate B-2

5.48 g (22.0 mmol) of Intermediate B-1, 5.31 g (22.0 mmol) of 1-bromo-4-chloronaphthalene, 1.27 g (1.1 mmol) of $Pd(PPh_3)_4$, and 4.5 g (33 mmol) of K_2CO_3 were dissolved in 200 mL of a THF/ H_2O (2/1 volume ratio) mixed solution, and then, the resulting solution was stirred at a temperature of $70^{\circ}C$ for 5 hours. After the obtained reaction solution was cooled to room temperature, 60 mL of water was added thereto, followed by three times of extraction using 60 mL of ethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and then, a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 4.68 g (yield: 75%) of Intermediate B-2. The prepared compound was identified by LC-MS.

($C_{16}H_{10}ClNO_2$: $M+$ 283.04)

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Synthesis of Intermediate B-3

2.83 g (10.00 mmol) of Intermediate B-2 and 8.3 g (50 mmol) of triethylphosphite were dissolved in 250 mL of cumene, and then, the resulting mixed solution was stirred under reflux in a nitrogen atmosphere for 24 hours. After the reaction was completed, an organic layer obtained by concentration under reduced pressure in high vacuum was separation-purified by silica gel column chromatography, thereby completing the preparation of 1.76 g (yield: 70%) of Intermediate B-3. The prepared compound was identified by LC-MS.

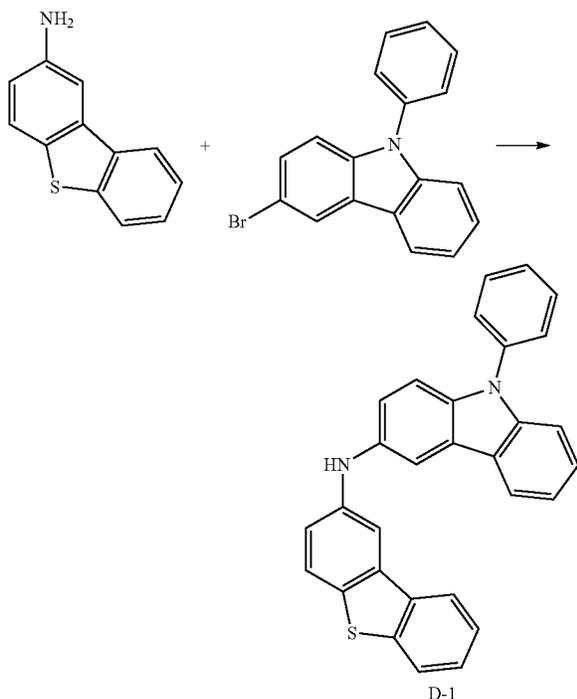
($C_{16}H_{10}ClN$: M+ 251.71)

Synthesis of Intermediate B-4

1.26 g (5.00 mmol) of Intermediate B-3, 0.95 g (6 mmol) of bromobenzene, 0.003 g (0.1 mmol) of $Pd(OAc)_2$, 0.1 g (0.5 mmol) of $P(tBu)_3$, and 0.84 g (7.5 mmol) of $KOtBu$ were dissolved in 100 mL of toluene, and then, the resulting mixed solution was stirred under reflux in a nitrogen atmosphere for 24 hours. After the reaction was completed, an extraction process was performed thereon three times using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 1.31 g (yield: 80%) of Intermediate B-4. The prepared compound was identified by LC-MS.

($C_{22}H_{14}ClN$: M+ 327.81)

Synthesis Example 10: Synthesis of Compound 59



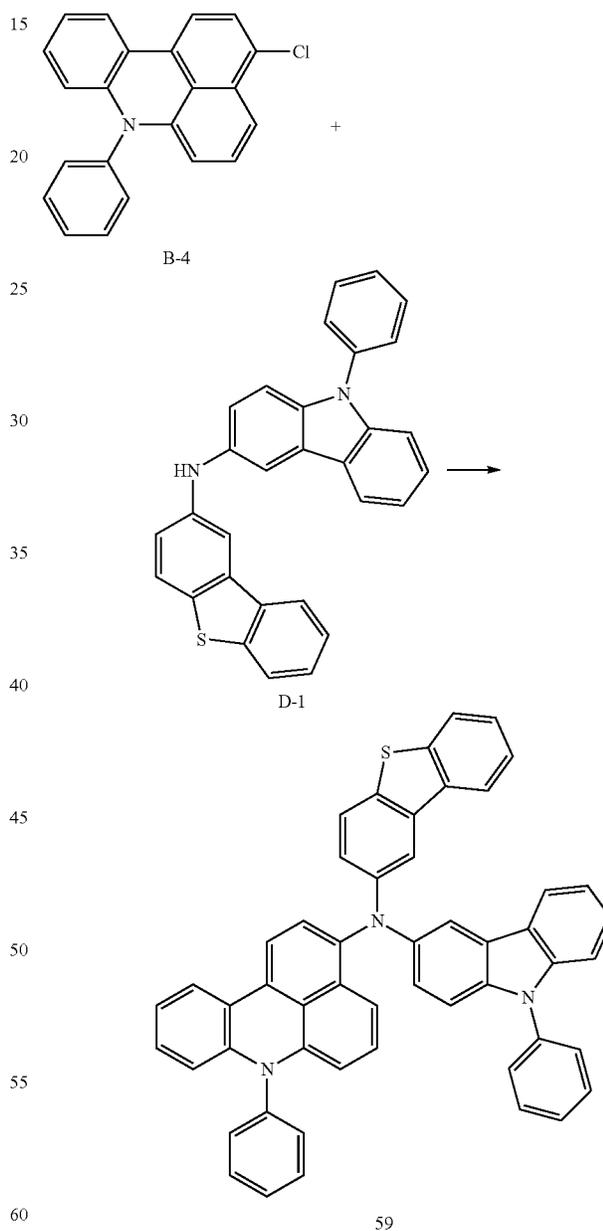
Synthesis of Intermediate D-1

5.98 g (30 mmol) of dibenzothiophene-2-amine, 6.44 g (20.0 mmol) of 3-bromo-9-phenyl-9H-carbazole, 0.37 g (0.4 mmol) of $Pd_2(dba)_3$, 0.08 g (0.4 mmol) of $P(tBu)_3$, and 5.76 g (60 mmol) of $KOtBu$ were dissolved in 200 mL of toluene,

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and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 7.40 g (yield: 84%) of Intermediate D-1. The prepared compound was identified by LC-MS.

($C_{30}H_{20}N_2S$: M+ 440.56)



Synthesis of Compound 59

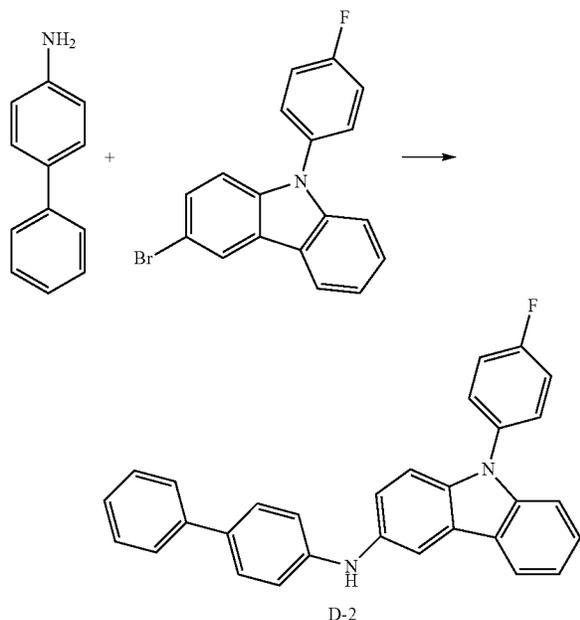
2.93 g (yield: 80%) of Compound 59 was obtained in the same (or substantially the same) manner as in Synthesis of Compound 10, except that 1.64 g (5 mmol) of Intermediate B-4 and 2.20 g (5 mmol) of Intermediate D-1 were used

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instead of Intermediate A-4 and Intermediate C-1, respectively. The prepared compound was identified by LC-MS and NMR.

($C_{52}H_{33}N_3S$: M+ 731.92)

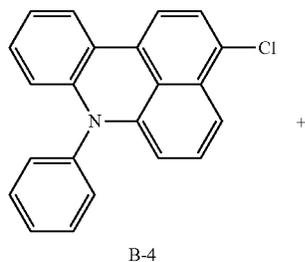
Synthesis Example 11: Synthesis of Compound 66



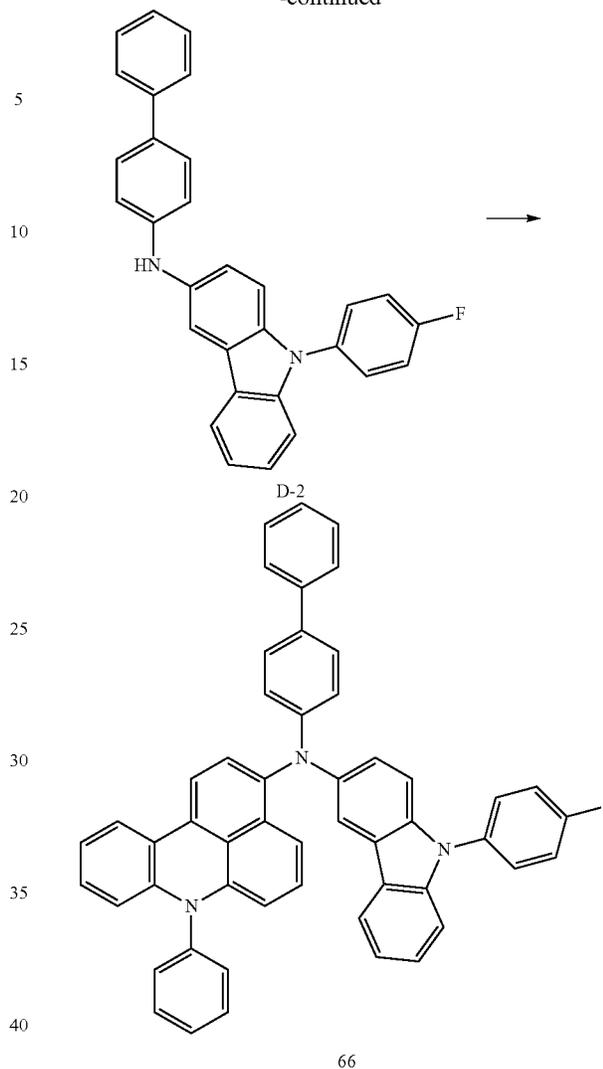
Synthesis of Intermediate D-2

5.08 g (30 mmol) of [1,1'-biphenyl]-4-amine, 6.80 g (20.0 mmol) of 3-bromo-9-(4-fluorophenyl)-9H-carbazole, 0.37 g (0.4 mmol) of $Pd_2(dba)_3$, 0.08 g (0.4 mmol) of $P(tBu)_3$, and 5.76 g (60 mmol) of $KOtBu$ were dissolved in 150 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon three times using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 7.28 g (yield: 85%) of Intermediate D-2. The prepared compound was identified by LC-MS.

($C_{30}H_{21}FN_2$: M+ 428.51)

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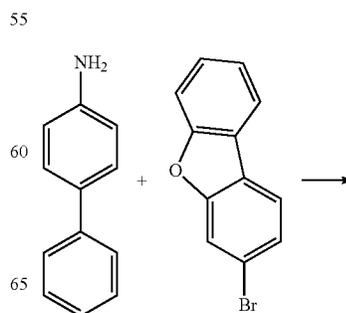
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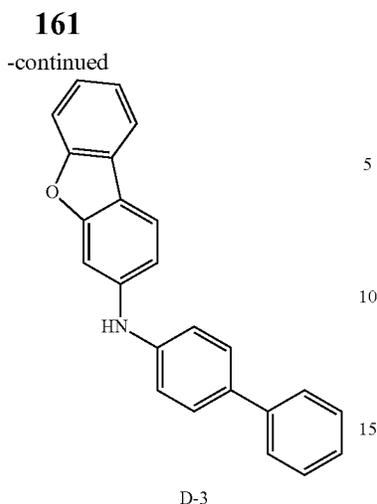


Synthesis of Compound 66

2.95 g (yield: 82%) of Compound 66 was obtained in the same (or substantially the same) manner as in Synthesis of Compound 10, except that 1.64 g (5 mmol) of Intermediate B-4 and 2.14 g (5 mmol) of Intermediate D-2 were used instead of Intermediate A-4 and Intermediate C-1, respectively. The prepared compound was identified by LC-MS and NMR. ($C_{52}H_{34}FN_3$: M+ 719.86)

Synthesis Example 12: Synthesis of Compound 69

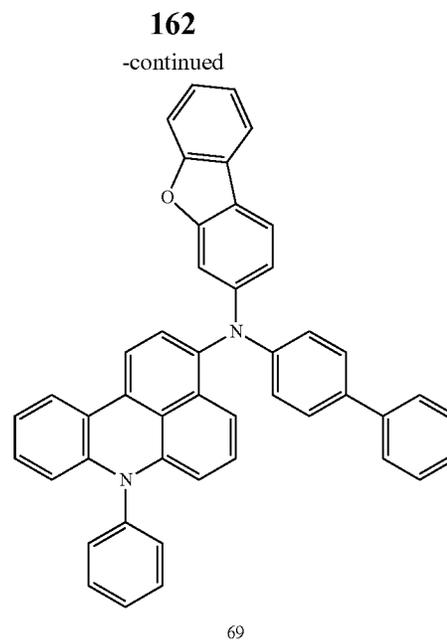
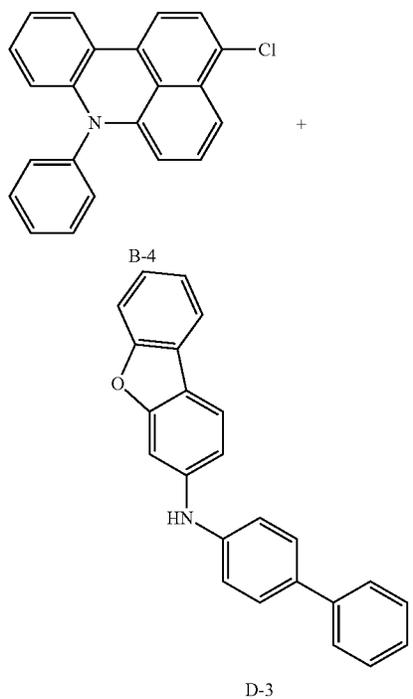




Synthesis of Intermediate D-3

5.08 g (30 mmol) of [1,1'-biphenyl]-4-amine, 4.94 g (20.0 mmol) of 3-bromodibenzo[b,d]furan, 0.37 g (0.4 mmol) of $\text{Pd}_2(\text{dba})_3$, 0.08 g (0.4 mmol) of $\text{P}(\text{tBu})_3$, and 5.76 g (60 mmol) of KOtBu were dissolved in 150 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon three times using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 5.84 g (yield: 87%) of Intermediate D-3. The prepared compound was identified by LC-MS.

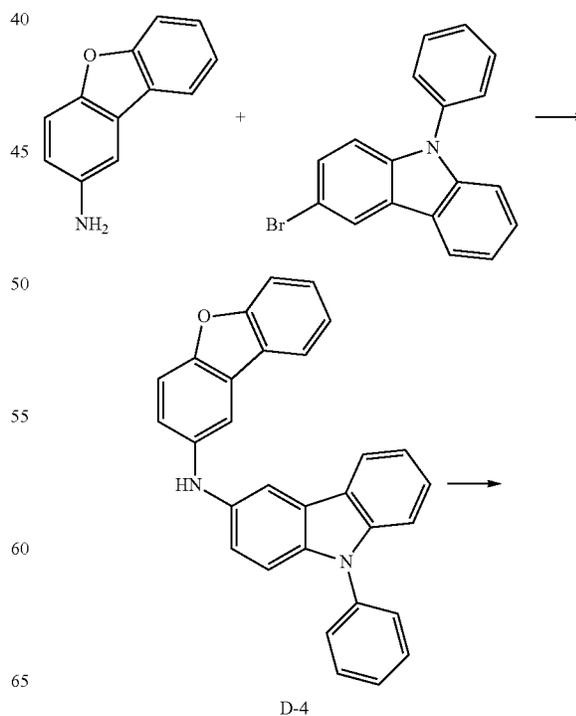
($\text{C}_{24}\text{H}_{17}\text{NO}$: M^+ 335.41)

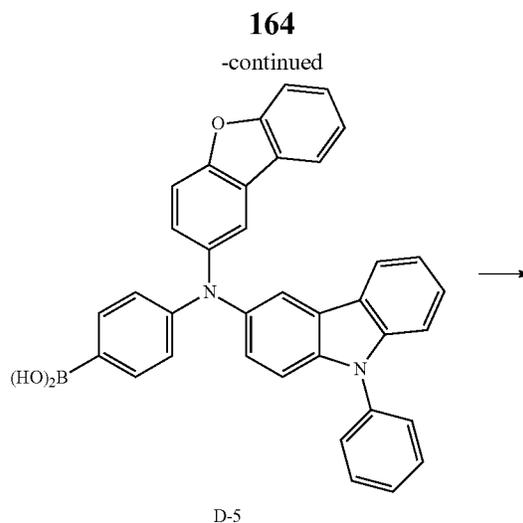
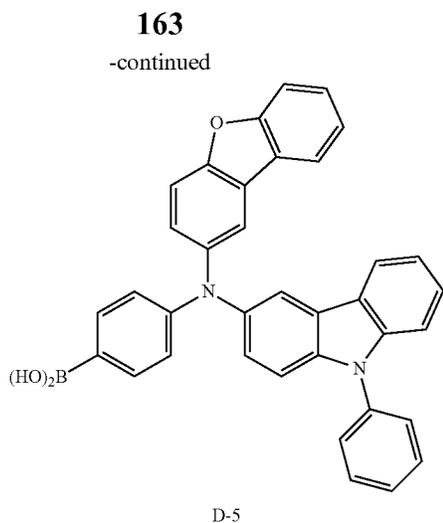


Synthesis of Compound 69

2.60 g (yield: 83%) of Compound 69 was obtained in the same (or substantially the same) manner as in Synthesis of Compound 10, except that 1.64 g (5 mmol) of Intermediate B-4 and 1.68 g (5 mmol) of Intermediate D-3 were used instead of Intermediate A-4 and Intermediate C-1, respectively. The prepared compound was identified by LC-MS and NMR. ($\text{C}_{46}\text{H}_{30}\text{N}_2\text{O}$: M^+ 626.76)

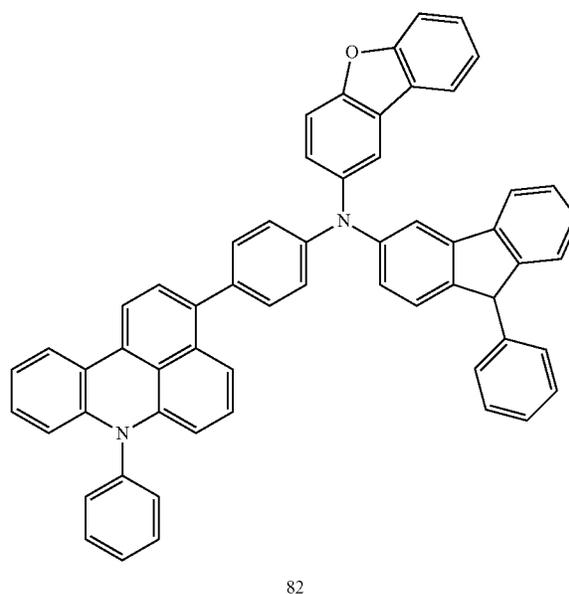
Synthesis Example 13: Synthesis of Compound 82





Synthesis of Intermediate D-4

5.50 g (30 mmol) of dibenzo[b,d]furan-2-amine, 6.44 g (20.0 mmol) of 3-bromo-9-phenyl-9H-carbazole, 0.37 g (0.4 mmol) of Pd₂(dba)₃, 0.08 g (0.4 mmol) of P(tBu)₃, and 5.76 g (60 mmol) of KOtBu were dissolved in 150 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon three times using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 7.13 g (yield: 84%) of Intermediate D-4. The prepared compound was identified by LC-MS. (C₃₀H₂₀N₂O: M+ 424.50)



Synthesis of Intermediate D-5

8.17 g (yield: 75%) of Intermediate D-5 was obtained in the same (or substantially the same) manner as in Synthesis of Intermediate C-6, except that 8.49 g (20 mmol) of Intermediate D-4 was used instead of Intermediate C-5. The prepared compound was identified by LC-MS.

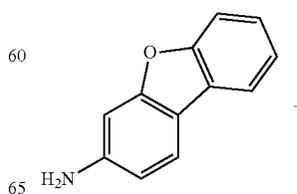
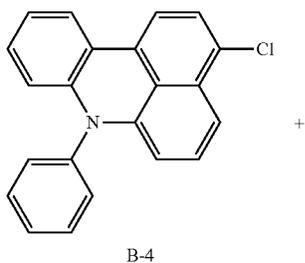
(C₃₆H₂₅BN₂O₃: M+ 544.42)

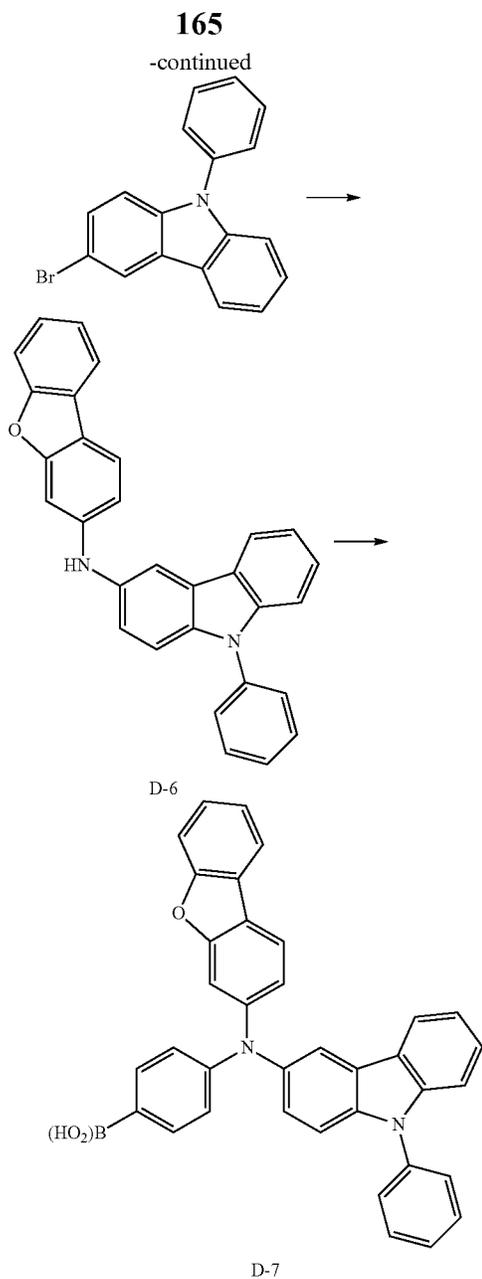
Synthesis of Compound 82

3.01 g (yield: 76%) of Compound 82 was obtained in the same (or substantially the same) manner as in Synthesis of Compound 29, except that 2.72 g (5 mmol) of Intermediate D-5 was used instead of Intermediate C-6, and Intermediate B-4 was used instead of Intermediate A-4. The prepared compound was identified by LC-MS and NMR.

(C₅₈H₃₇N₃O: M+ 791.95)

Synthesis Example 14: Synthesis of Compound 84





Synthesis of Intermediate D-6

5.50 g (30 mmol) of dibenzo[b,d]furan-3-amine, 6.44 g (20.0 mmol) of 3-bromo-9-phenyl-9H-carbazole, 0.37 g (0.4 mmol) of $\text{Pd}_2(\text{dba})_3$, 0.08 g (0.4 mmol) of $\text{P}(\text{tBu})_3$, and 5.76 g (60 mmol) of KOtBu were dissolved in 150 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon three times using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 6.79 g (yield: 80%) of Intermediate D-6. The prepared compound was identified by LC-MS.

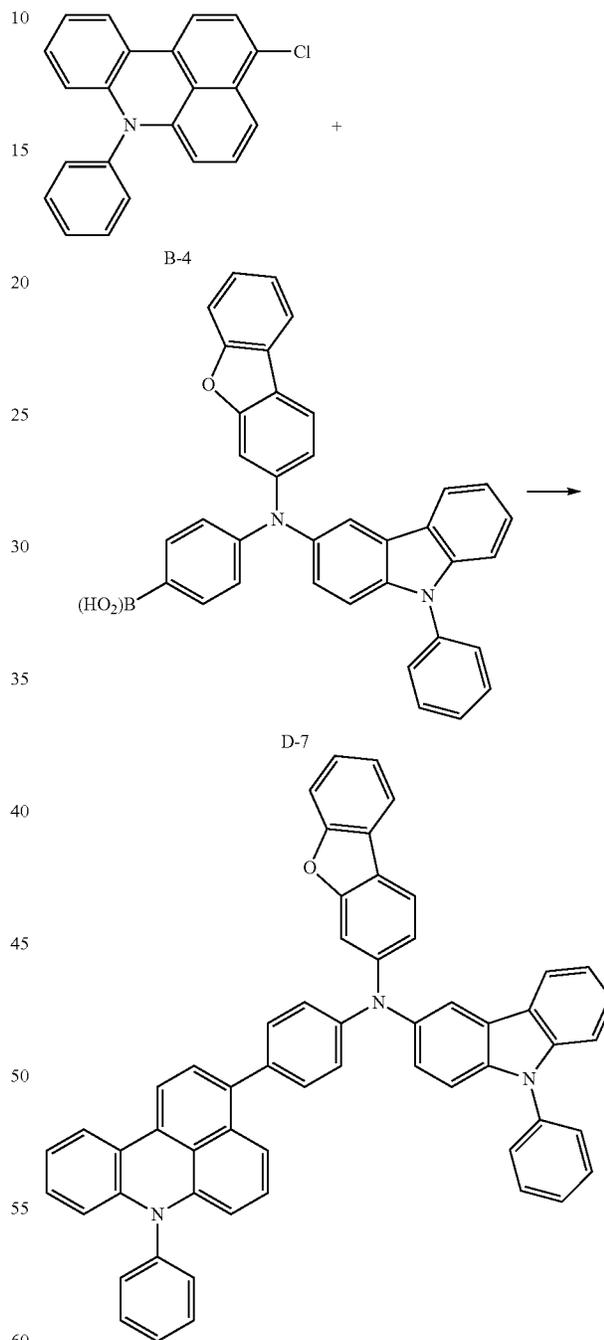
($\text{C}_{30}\text{H}_{20}\text{N}_2\text{O}$: M^+ 424.50)

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Synthesis of Intermediate D-7

7.73 g (yield: 71%) of Intermediate D-7 was obtained in the same (or substantially the same) manner as in Synthesis of Intermediate C-6, except that 8.49 g (20 mmol) of Intermediate D-6 was used instead of Intermediate C-5. The prepared compound was identified by LC-MS.

($\text{C}_{36}\text{H}_{25}\text{BN}_2\text{O}_3$: M^+ 544.42)



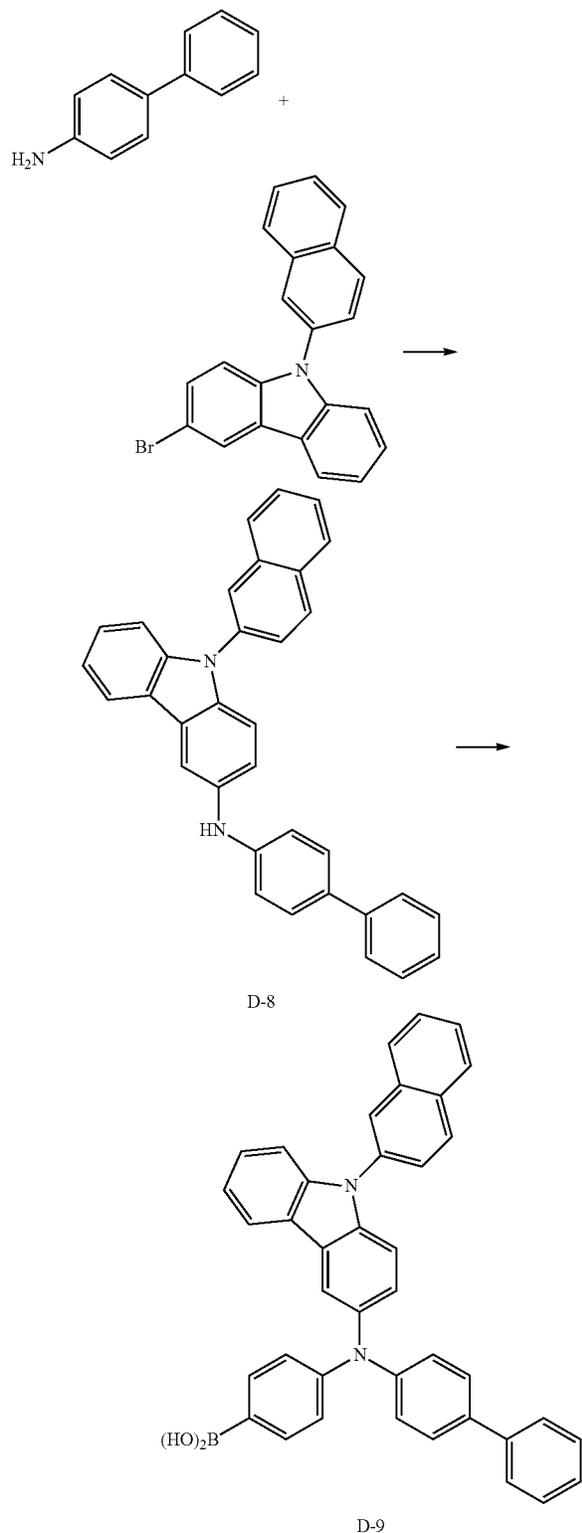
Synthesis of Compound 84

2.93 g (yield: 74%) of Compound 84 was obtained in the same (or substantially the same) manner as in Synthesis of Compound 29, except that 2.72 g (5 mmol) of Intermediate

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D-7 was used instead of Intermediate C-6. The prepared compound was identified by LC-MS and NMR. ($C_{58}H_{37}N_3O$: M+ 791.95)

Synthesis Example 15: Synthesis of Compound 89



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Synthesis of Intermediate D-8

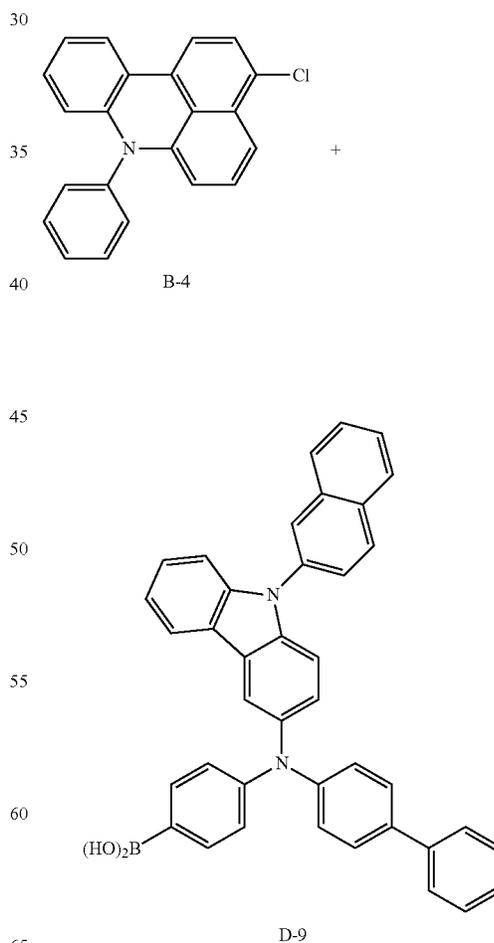
5.08 g (30 mmol) of [1,1'-biphenyl]-4-amine, 7.45 g (20.0 mmol) of 3-bromo-9-(naphthalen-2-yl)-9H-carbazole, 0.37 g (0.4 mmol) of $Pd_2(dba)_3$, 0.08 g (0.4 mmol) of $P(tBu)_3$, and 5.76 g (60 mmol) of $KOtBu$ were dissolved in 150 mL of toluene, and then, the resulting mixed solution was stirred at a temperature of 85° C. for 4 hours. After the obtained reaction solution was cooled to room temperature, an extraction process was performed thereon three times using 50 mL of water and 50 mL of diethylether. An organic layer obtained from the resulting solution was dried using magnesium sulfate, and a solvent was removed therefrom by evaporation. The residue obtained therefrom was separation-purified by silica gel column chromatography, thereby completing the preparation of 6.91 g (yield: 75%) of Intermediate D-8. The prepared compound was identified by LC-MS.

($C_{34}H_{24}N_2$: M+ 460.58)

Synthesis of Intermediate D-9

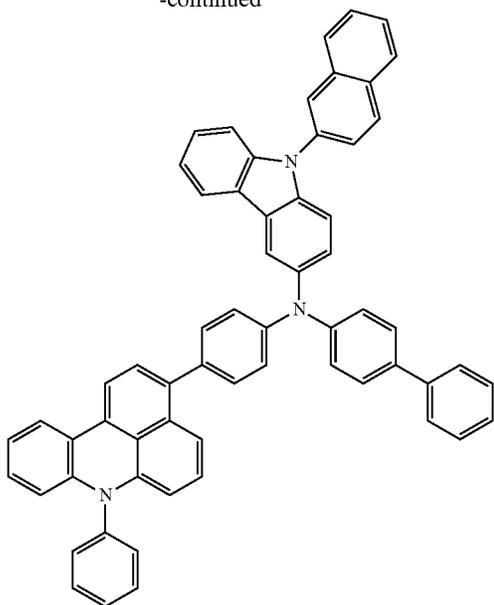
8.48 g (yield: 73%) of Intermediate D-9 was obtained in the same (or substantially the same) manner as in Synthesis of Intermediate C-6, except that 9.21 g (20 mmol) of Intermediate D-8 was used instead of Intermediate C-5. The prepared compound was identified by LC-MS.

($C_{40}H_{29}BN_2O_2$: M+ 580.49)



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-continued



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Synthesis of Compound 89

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3.23 g (yield: 78%) of Compound 89 was obtained in the same (or substantially the same) manner as in Synthesis of Compound 29, except that 2.90 g (5 mmol) of Intermediate D-9 was used instead of Intermediate C-6, and Intermediate B-4 was used instead of Intermediate A-4. The prepared compound was identified by LC-MS and NMR. ($C_{62}H_{41}N_3$: M^+ 828.03)

The compounds synthesized according to Synthesis Examples 1 to 15 were identified by 1H NMR and mass spectrometry/fast atom bombardment (MS/FAB), and results are shown in Table 1 below.

TABLE 1

Compound	1H NMR ($CDCl_3$, 400 MHz)	MS/FAB	
		found	calc.
10	8.21 (d, 1H), 8.03 (d, 1H), 7.82 (1H, d), 7.67 (dd, 2H), 7.67-7.49 (m, 7H), 7.40-7.36 (m, 6H), 7.30-7.24 (m, 5H), 7.11-7.07 (m, 3H), 6.94 (d, 1h), 6.62-6.57 (m, 2H), 6.40-6.37 (m, 2H), 6.19 (dd, 2H)	715.86	715.26
14	8.21 (d, 1H), 8.03 (d, 1H), 7.85 (d, 1H), 7.67 (dd, 2H), 7.57-7.49 (m, 5H), 7.44-7.29 (m, 5H), 7.27-7.07 (m, 19H), 6.96 (d, 1H), 6.80 (d, 1H), 6.67-6.61 (m, 3H), 6.35-6.31 (m, 2H), 6.19 (dd, 2H)	866.08	865.35
18	8.21 (d, 1H), 8.03 (d, 1H), 7.85 (d, 1H), 7.67 (dd, 2H), 7.57 (d, 1H), 7.50-7.48 (m, 4H), 7.44-7.29 (m, 5H), 7.27-7.23 (m, 10H), 7.23-7.07 (m, 9H), 6.96 (d, 1H), 6.80 (d, 1H), 6.67-6.61 (m, 3H), 6.35-6.31 (m, 2H), 6.19 (dd, 2H)	719.86	719.27
22	8.03 (d, 1H), 7.76 (d, 1H), 7.67 (dd, 2H), 7.58-7.56 (m, 3H), 7.46-7.45 (m, 5H), 7.37-7.24 (m, 4H), 7.11-7.07 (m, 5H), 6.67-6.73 (m, 2H), 6.62-6.35 (m, 5H), 6.19 (dd, 2H), 1.61 (m, 6H)	652.84	652.29
29	8.21 (d, 1H), 7.92 (d, 1H), 7.80-7.77 (m, 3H), 7.67-7.57 (m, 7H), 7.50-7.24 (m, 20H), 7.15-7.07 (m, 3H), 6.84 (d, 2H), 6.70-6.48 (m, 6H)	854.07	853.35
36	8.21 (d, 1H), 7.92 (d, 1H), 7.81-7.79 (m, 4H), 7.67 (dd, 2H), 7.57-7.49 (m, 6H), 7.43-7.36 (m, 7H), 7.30-7.23 (m, 5H), 7.15-7.03 (m, 5H), 6.79 (d, 2H), 6.62-6.57 (m, 2H), 6.48 (dd, 2H)	791.95	791.29
41	8.25 (d, 1H), 7.92-7.87 (m, 3H), 7.80-7.79 (d, 2H), 7.71-7.67 (m, 3H), 7.57-7.51 (m, 6H), 7.46-7.42 (m, 8H), 7.38-7.37 (m, 4H), 7.27-7.25 (m, 3H), 7.15-7.07 (m, 3H), 6.66-6.59 (m, 6H), 6.48 (dd, 2H)	828.03	827.33
48	7.92-7.88 (m, 4H), 7.80-7.79 (d, 2H), 7.67-7.65 (d, 2H), 7.57-7.54 (m, 3H), 7.46-7.42 (m, 10H), 7.37-7.15 (m, 7H), 7.07-7.05 (m, 2H), 6.85-6.74 (m, 4H), 6.62-6.46 (m, 8H)	851.07	850.33
54	8.21 (d, 1H), 7.99-7.97 (d, 2H), 7.86-7.76 (d, 2H), 7.68-7.50 (m, 7H), 7.49-7.37 (m, 8H), 7.36-7.04 (m, 10H), 7.01-6.62 (m, 4H), 6.52-6.50 (m, 2H), 6.40-6.38 (d, 1H)	751.93	751.30

Compound	¹ H NMR (CDCl ₃ , 400 MHz)	MS/FAB	
		found	calc.
59	8.21 (d, 1H), 8.05 (d, 1H), 7.88-7.70 (m, 4H), 7.67-7.60 (m, 2H), 7.53-7.44 (m, 6H), 7.41-7.22 (m, 7H), 7.13-7.01 (m, 6H), 6.85-6.81 (m, 2H), 6.62-6.60 (m, 2H), 6.52-6.50 (m, 2H)	731.92	731.24
66	8.21 (d, 1H), 7.88 (d, 1H), 7.76 (d, 1H), 7.67 (d, 1H), 7.57-7.55 (m, 2H), 7.46-7.22 (m, 13H), 7.13-7.01 (m, 8H), 6.91-6.88 (m, 2H), 6.62-6.52 (m, 5H)	719.86	719.27
69	7.88 (d, 1H), 7.81 (d, 1H), 7.76 (d, 2H), 7.67 (d, 1H), 7.57-7.53 (m, 3H), 7.46-7.43 (m, 4H), 7.38-7.36 (m, 2H), 7.23-7.21 (m, 2H), 7.13-7.01 (m, 8H), 6.91 (d, 1H), 6.65-6.62 (m, 3H), 6.52-6.50 (m, 2H)	626.76	626.24
82	8.21 (d, 1H), 8.09 (d, 1H), 7.82-7.80 (d, 2H), 7.63 (d, 1H), 7.53-7.49 (m, 8H), 7.40-7.00 (m, 18H), 6.62-6.60 (m, 4H), 6.52-6.50 (m, 2H)	791.95	791.29
84	8.21 (d, 1H), 8.09 (d, 1H), 7.81-7.79 (m, 3H), 7.63 (d, 1H), 7.53-7.48 (m, 7H), 7.39-7.23 (m, 10H), 7.16-7.01 (m, 8H), 6.73-6.71 (m, 2H), 6.62-6.52 (m, 4H)	791.95	791.29
89	8.25 (d, 1H), 8.09 (d, 1H), 7.79-7.87 (m, 2H), 7.80-7.71 (m, 2H), 7.63-7.25 (m, 21H), 7.16-7.02 (m, 6H), 6.65-6.52 (m, 8H)	828.03	827.33

Example 1

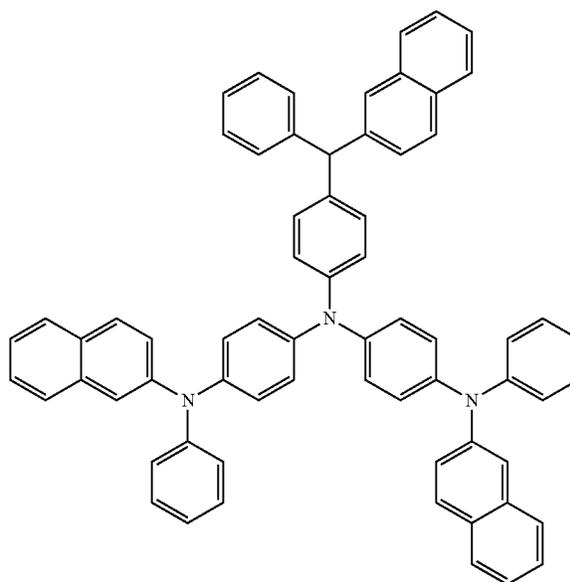
An anode was prepared by cutting an indium tin oxide (ITO) glass substrate (manufactured by Corning Inc.) on which ITO was formed to a thickness of 15 Ω/cm² (1,200 Å) to a size of 50 mm×50 mm×0.7 mm, ultrasonically cleaning the ITO glass substrate using isopropyl alcohol and pure water, for 5 minutes in each, and then, exposing to irradiation of UV light for 30 minutes and ozone to clean. Then, the obtained ITO glass substrate was loaded into a vacuum deposition apparatus.

2-TNATA was vacuum deposited on the ITO glass substrate to form a hole injection layer having a thickness of 600 Å, and then, Compound 10 was vacuum deposited on the hole injection layer to form a hole transport layer having a thickness of 300 Å, thereby forming a hole transport region.

9,10-di-naphthalene-2-yl-anthracene (ADN) and N,N,N',N'-tetraphenyl-pyrene-1,6-diamine (TPD) were co-deposited on the hole transport region at a weight ratio of 98:2 to form an emission layer having a thickness of 300 Å.

Alq₃ was deposited on the emission layer to form an electron transport layer having a thickness of 300 Å, LiF was deposited on the electron transport layer to form an electron injection layer having a thickness of 10 Å, and Al was vacuum deposited on the electron injection layer to form a

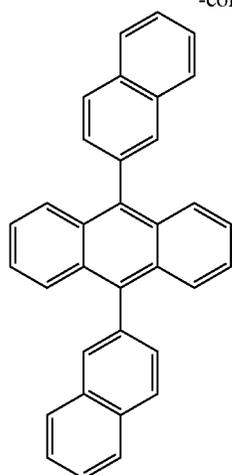
cathode (i.e., a cathode electrode) having a thickness of 3,000 Å, thereby completing the manufacture of an organic light-emitting device.



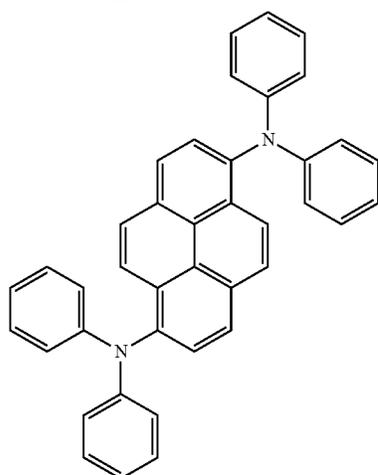
2-TNATA

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ADN



TPD

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Examples 2 to 15 and Comparative Examples 1 to 3

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Organic light-emitting devices were each manufactured in
 10 the same (or substantially the same) manner as in Synthesis of Example 1, except that Compounds 14, 18, 22, 29, 36, 41, 48, 54, 59, 66, 69, 82, 84, and 89, NPB, Compound A, and
 15 Compound B were used instead of Compound 10 in forming the hole transport layer.

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Evaluation Example 1

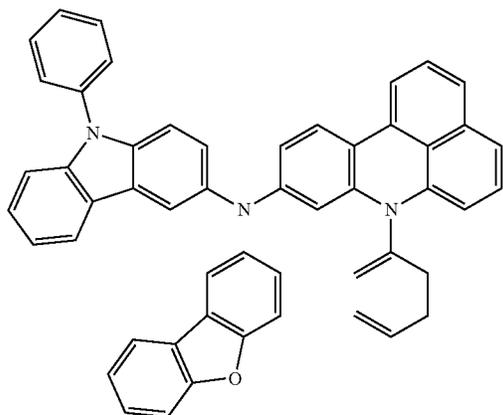
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30 The driving voltage, current density, brightness, efficiency, and half-lifespan of the organic light-emitting devices of Examples 1 to 15 and Comparative Examples 1 to 3 were evaluated by using a Keithley SMU 236 meter and
 35 a PR650 brightness measuring meter. Results thereof are shown in Table 2. Here, the half-lifespan results were obtained by measuring the time at which the brightness of
 40 the organic light-emitting device reached 50% of the initial brightness after being driven.

TABLE 2

	Material for forming a hole transport layer	Driving voltage (V)	Current density (mA/cm ²)	Brightness (cd/cm ²)	Efficiency (cd/A)	Emission color	Half-life span (hr @100 mA/cm ²)
Example 1	Compound 10	5.71	50	3040	6.08	Blue	345
Example 2	Compound 14	5.65	50	3085	6.17	Blue	355
Example 3	Compound 18	5.54	50	3125	6.25	Blue	334
Example 4	Compound 22	5.50	50	3155	6.31	Blue	340
Example 5	Compound 29	5.48	50	3110	6.22	Blue	325
Example 6	Compound 36	5.39	50	3210	6.42	Blue	351
Example 7	Compound 41	5.43	50	3170	6.34	Blue	338
Example 8	Compound 48	5.53	50	3135	6.27	Blue	375
Example 9	Compound 54	5.70	50	3060	6.12	Blue	348
Example 10	Compound 59	5.68	50	3095	6.19	Blue	342
Example 11	Compound 66	5.65	50	3120	6.24	Blue	352
Example 12	Compound 69	5.60	50	3190	6.38	Blue	364
Example 13	Compound 82	5.40	50	3275	6.55	Blue	365
Example 14	Compound 84	5.42	50	3235	6.47	Blue	378
Example 15	Compound 89	5.51	50	3210	6.42	Blue	342
Comparative Example 1	NPB	7.01	50	2645	5.29	Blue	258
Comparative Example 2	Compound A	7.24	50	2550	5.10	Blue	243
Comparative Example 3	Compound B	6.28	50	3020	6.04	Blue	284

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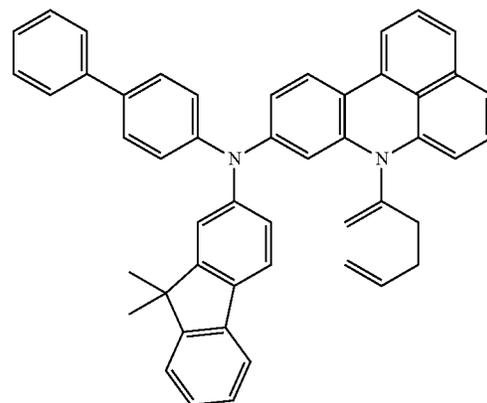
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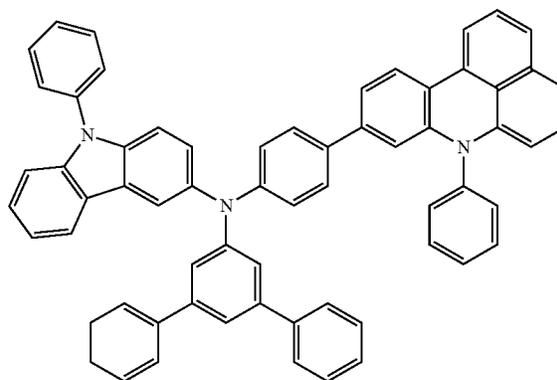
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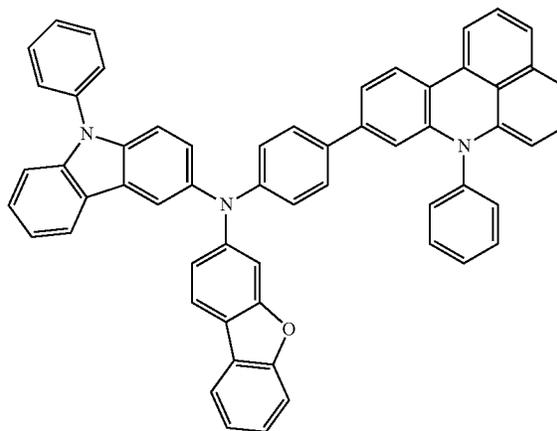
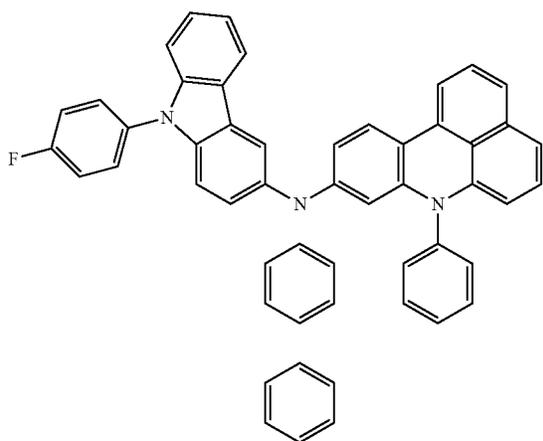
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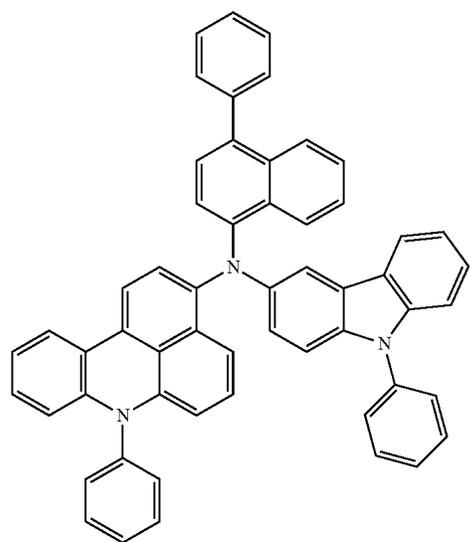
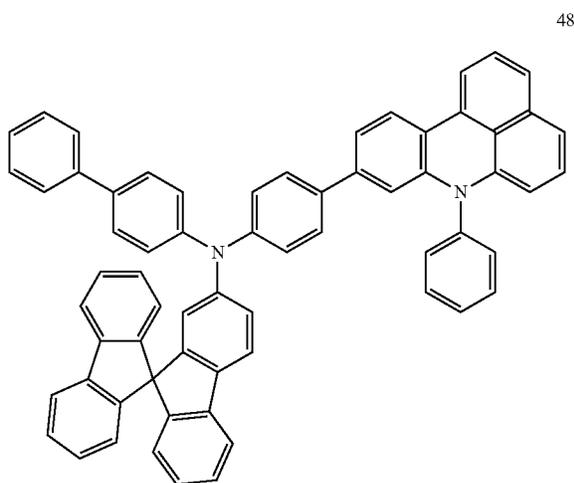
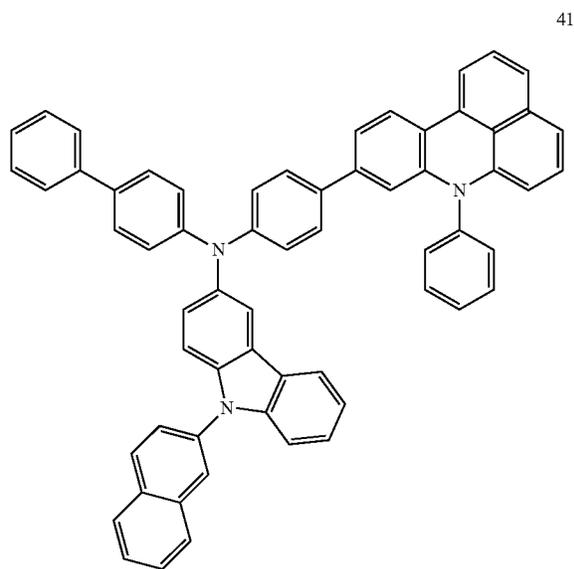
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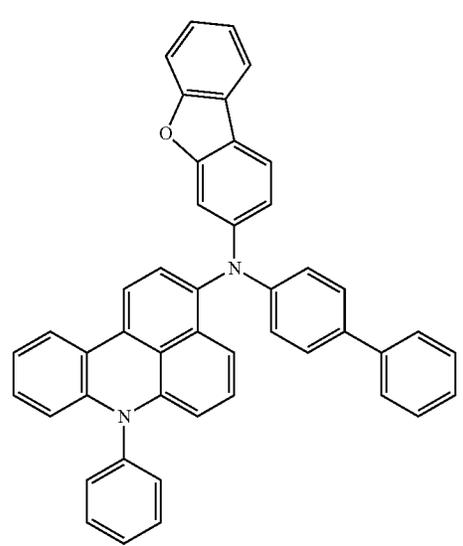
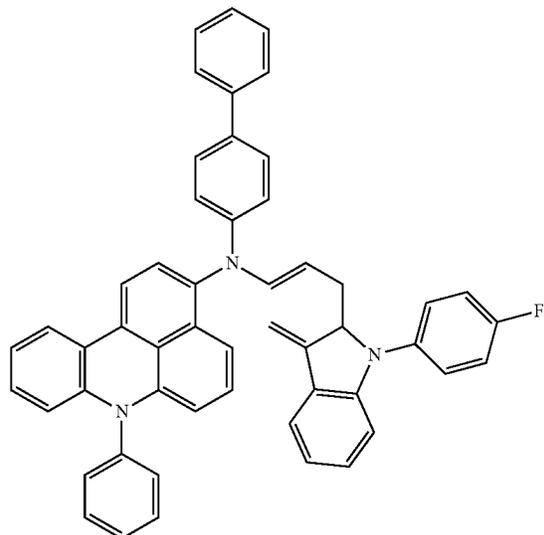
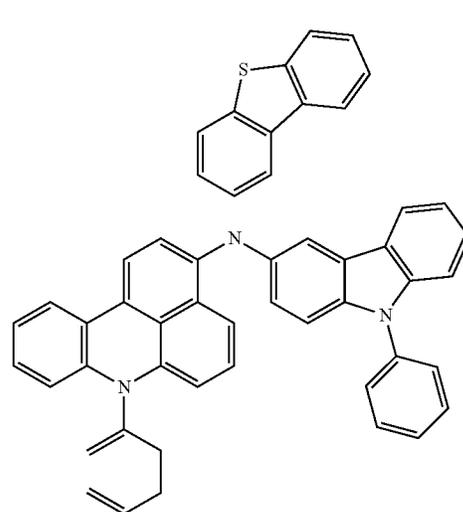
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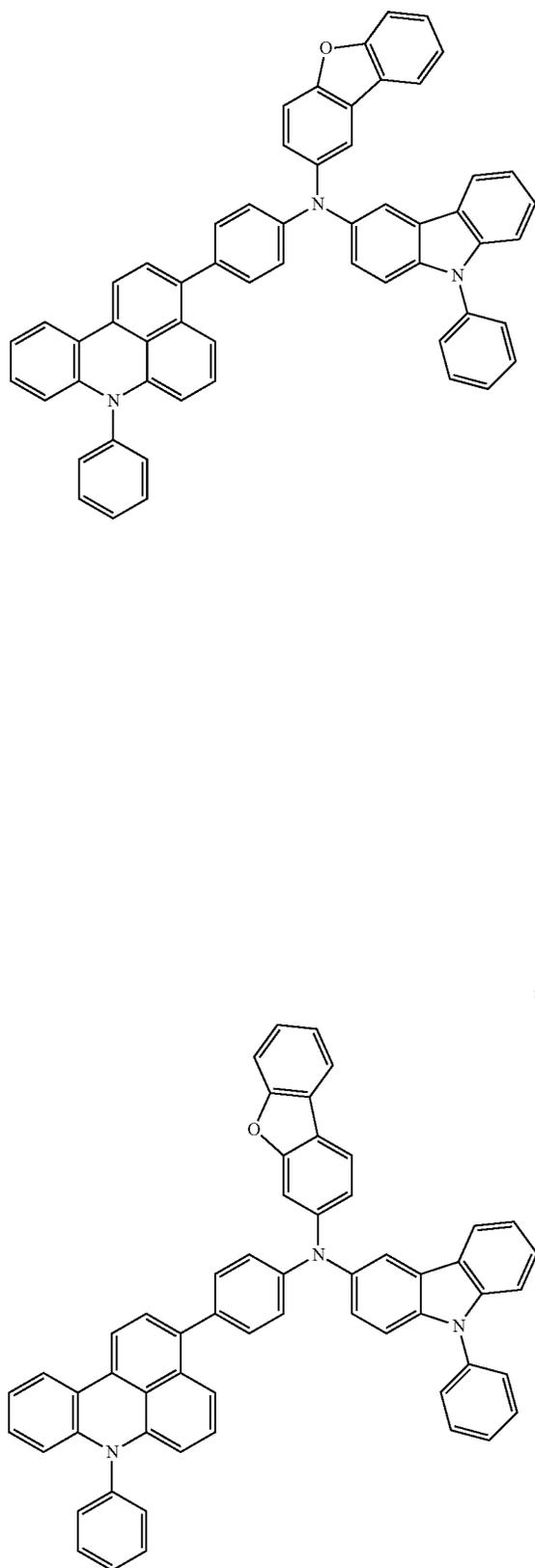
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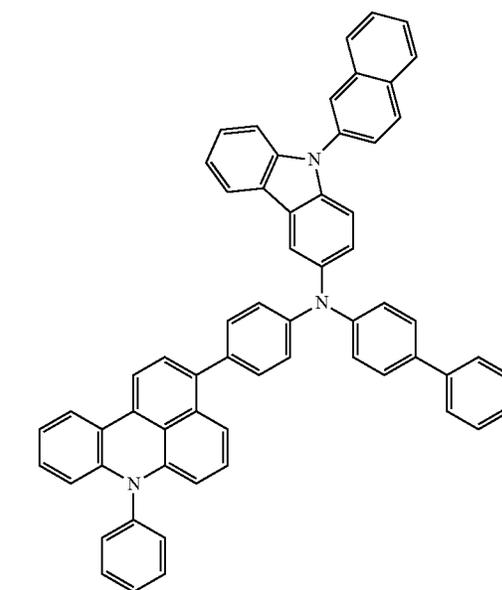
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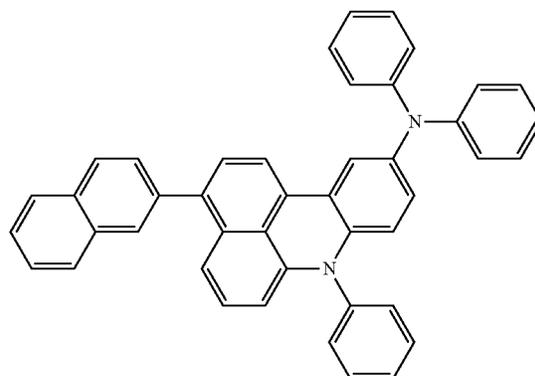
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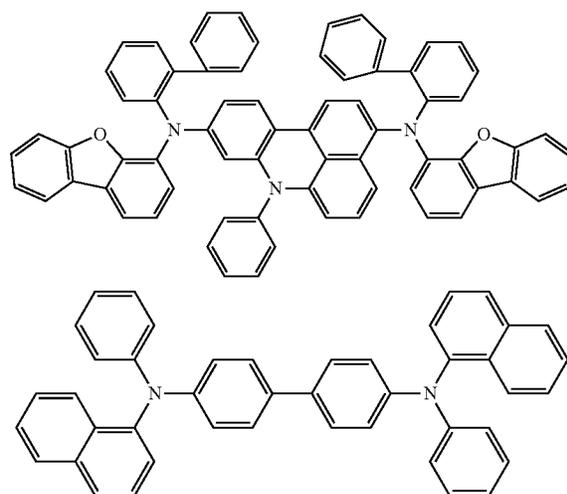
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Compound A



Compound B



NPB

65 Referring to the results shown in Table 2, it can be seen that the organic light-emitting devices of Examples 1 to 15 had low driving voltage and improved brightness, efficiency,

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and half-lifespans, as compared to those of the organic light-emitting devices of Comparative Examples 1 to 3.

As used herein, the terms “use,” “using,” and “used” may be considered synonymous with the terms “utilize,” “utilizing,” and “utilized,” respectively.

In addition, the terms “substantially,” “about,” and similar terms are used as terms of approximation and not as terms of degree, and are intended to account for the inherent deviations in measured or calculated values that would be recognized by those of ordinary skill in the art.

Also, any numerical range recited herein is intended to include all sub-ranges of the same numerical precision subsumed within the recited range. For example, a range of “1.0 to 10.0” is intended to include all subranges between (and including) the recited minimum value of 1.0 and the recited maximum value of 10.0, that is, having a minimum value equal to or greater than 1.0 and a maximum value equal to or less than 10.0, such as, for example, 2.4 to 7.6. Any maximum numerical limitation recited herein is intended to include all lower numerical limitations subsumed therein and any minimum numerical limitation recited in this specification is intended to include all higher numerical limitations subsumed therein. Accordingly, Applicant reserves the right to amend this specification, including the claims, to expressly recite any sub-range subsumed within the ranges expressly recited herein.

It should be understood that example embodiments described herein should be considered in a descriptive sense only and not for purposes of limitation. Descriptions of features or aspects within each example embodiment should typically be considered as available for other similar features or aspects in other example embodiments.

The novel condensed cyclic compound according to embodiments of the present disclosure may have excellent hole injection characteristics. Accordingly, the novel condensed cyclic compound is suitable for use as a hole transport material for red, green, and/or white fluorescent and/or phosphorescent devices. An organic light-emitting device including the condensed cyclic compound according to the present embodiments may have high efficiency, low driving voltage, high luminance, and a long lifespan.

It should be understood that embodiments described herein should be considered in a descriptive sense only and not for purposes of limitation. Descriptions of features or aspects within each embodiment should typically be considered as available for other similar features or aspects in other embodiments.

While one or more embodiments have been described with reference to the drawing, it will be understood by those of ordinary skill in the art that various changes in form and details may be made therein without departing from the spirit and scope of the present disclosure as defined by the following claims and equivalents thereof.

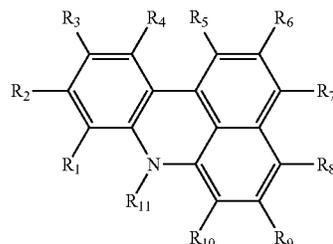
What is claimed is:

1. An organic light-emitting device comprising:
 - a first electrode;
 - a second electrode facing the first electrode; and
 - an organic layer between the first electrode and the second electrode, the organic layer comprising:
 - i) an emission layer,
 - ii) a hole transport region between the first electrode and the emission layer, the hole transport region comprising at least one selected from a hole injection layer, a hole transport layer, a buffer layer, and an electron blocking layer; and
 - iii) an electron transport region between the emission layer and the second electrode, the electron transport

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region comprising at least one selected from a hole blocking layer, an electron transport layer, and an electron injection layer,

wherein the emission layer is to emit blue fluorescence, wherein the hole transport region comprises a condensed cyclic compound represented by Formula 1:

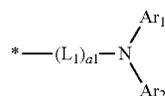


Formula 1

wherein, in Formula 1,

R_1 to R_6 and R_8 to R_{11} are each independently selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a substituted or unsubstituted C_1 - C_{60} alkyl group, a substituted or unsubstituted C_2 - C_{60} alkenyl group, a substituted or unsubstituted C_2 - C_{60} alkynyl group, a substituted or unsubstituted C_1 - C_{60} alkoxy group, a substituted or unsubstituted C_3 - C_{10} cycloalkyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkyl group, a substituted or unsubstituted C_3 - C_{10} cycloalkenyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkenyl group, a substituted or unsubstituted C_6 - C_{60} aryl group, a substituted or unsubstituted C_6 - C_{60} aryloxy group, a substituted or unsubstituted C_6 - C_{60} arylthio group, a substituted or unsubstituted C_1 - C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, and —Si(Q_1)(Q_2)(Q_3), and

R_7 is a group represented by Formula 2:



Formula 2

wherein in Formula 2,

L_1 is selected from a substituted or unsubstituted C_3 - C_{10} cycloalkylene group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkylene group, a substituted or unsubstituted C_3 - C_{10} cycloalkenylene group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkenylene group, a substituted or unsubstituted C_6 - C_{60} arylene group, a substituted or unsubstituted C_1 - C_{60} heteroarylene group, a substituted or unsubstituted divalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted divalent non-aromatic condensed heteropolycyclic group;

a_1 is an integer selected from 0 to 3, wherein when a_1 is two or more, two or more L_1 (s) are identical to or different from each other; and

Ar₁ and Ar₂ are each independently selected from a substituted or unsubstituted fluorenyl group, a substituted or unsubstituted spiro-bifluorenyl group, a substituted or unsubstituted carbazolyl group, a substituted or unsubstituted dibenzofuranyl group, a substituted or unsubstituted dibenzothiophenyl group, and a substituted or unsubstituted dibenzosilolyl group,

provided that Ar₁ and Ar₂ are different from each other, and

provided that at least one selected from Ar₁ and Ar₂ is a carbazolyl group substituted with a C₆-C₆₀ aryl group substituted with at least one selected from —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, and a phosphoric acid group or a salt thereof,

wherein at least one substituent of the substituted fluorenyl group, substituted spiro-bifluorenyl group, substituted carbazolyl group, substituted dibenzofuranyl group, substituted dibenzothiophenyl group, substituted dibenzosilolyl group, substituted C₃-C₁₀ cycloalkylene group, substituted C₁-C₁₀ heterocycloalkylene group, substituted C₃-C₁₀ cycloalkenylene group, substituted C₁-C₁₀ heterocycloalkenylene group, substituted C₆-C₆₀ arylene group, substituted C₁-C₆₀ heteroarylene group, substituted divalent non-aromatic condensed polycyclic group, substituted divalent non-aromatic condensed heteropolycyclic group, substituted C₁-C₆₀ alkyl group, substituted C₂-C₆₀ alkenyl group, substituted C₂-C₆₀ alkynyl group, substituted C₁-C₆₀ alkoxy group, substituted C₃-C₁₀ cycloalkyl group, substituted C₁-C₁₀ heterocycloalkyl group, substituted C₃-C₁₀ cycloalkenyl group, substituted C₁-C₁₀ heterocycloalkenyl group, substituted C₆-C₆₀ aryl group, substituted C₆-C₆₀ aryloxy group, substituted C₆-C₆₀ arylthio group, substituted C₁-C₆₀ heteroaryl group, substituted monovalent non-aromatic condensed polycyclic group, and substituted monovalent non-aromatic condensed heteropolycyclic group is selected from the group consisting of:

deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group,

a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, and —Si(Q₁₁)(Q₁₂)(Q₁₃),

a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀

aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, a biphenyl group, and a terphenyl group, and

a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, a biphenyl group, and a terphenyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, a biphenyl group, a terphenyl group, and —Si(Q₂₁)(Q₂₂)(Q₂₃); and —Si(Q₃₁)(Q₃₂)(Q₃₃),

wherein Q₁ to Q₃, Q₁₁ to Q₁₃, Q₂₁ to Q₂₃, and Q₃₁ to Q₃₃ are each independently selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, a biphenyl group, and a terphenyl group.

2. The organic light-emitting device of claim 1, wherein L₁ is selected from the group consisting of:

a phenylene group, a pentalenylene group, an indenylene group, a naphthylene group, an azulenylene group, a heptalenylene group, an indacenylene group, an acenaphthylene group, a fluorenylene group, spiro-bifluorenylene group, a benzofluorenylene group, a dibenzofluorenylene group, a phenalenylene group, a phenanthrenylene group, an anthracenylene group, a fluoranthenylene group, a triphenylenylene group, a pyrenylene group, a chrysenylene group, a naphthacenylenylene group, a picenylene group, a perylenylene group, a pentaphenylene group, a hexacenylene group, a pentacenylene group, a rubicenylene group, a coronenylene group, an ovalenylene group, a pyrrolylene group, a thiophenylene group, a furanylene group, an imidazolylene group, a pyrazolylene group, a thiazolylene group, an isothiazolylene group, an oxazolylene group, an isoxazolylene group, a pyridinylene group, a pyrazinylene group, a pyrimidinylene group, a pyridazinylene group, an isoindolylene group, an indolylene group, an indazolylene group, a purinylene

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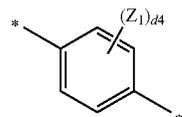
group, a quinolinylene group, an isoquinolinylene group, a benzoquinolinylene group, a phthalazinylene group, a naphthyridinylene group, a quinoxalinylene group, a quinazolinylene group, a cinnolinylene group, a carbazolylene group, a phenanthridinylene group, an acridinylene group, a phenanthrolinylene group, a phenazinylene group, a benzoimidazolylene group, a benzofuranylene group, a benzothiophenylene group, an isobenzothiazolylene group, a benzoxazolylene group, an isobenzoxazolylene group, a triazolylene group, a tetrazolylene group, an oxadiazolylene group, a triazinylene group, a dibenzofuranylene group, a dibenzothiophenylene group, a dibenzosilolylene group, a benzocarbazolylene group, a dibenzocarbazolylene group, a thiadiazolylene group, an imidazopyridinylene group, and an imidazopyrimidinylene group, and

a phenylene group, a pentalenylene group, an indenylene group, a naphthylene group, an azulenylene group, a heptalenylene group, an indacenylene group, an acenaphthylene group, a fluorenylene group, spiro-bifluorenylene group, a benzofluorenylene group, a dibenzofluorenylene group, a phenalenylene group, a phenanthrenylene group, an anthracenylene group, a fluoranthenylene group, a triphenylenylene group, a pyrenylene group, a chrysenylene group, a naphthacenylenylene group, a picenylene group, a perylenylene group, a pentaphenylene group, a hexacenylene group, a pentacenylene group, a rubicenylene group, a coronenylene group, an ovalenylene group, a pyrrolylene group, a thiophenylene group, a furanylene group, an imidazolylene group, a pyrazolylene group, a thiazolylene group, an isothiazolylene group, an oxazolylene group, an isoxazolylene group, a pyridinylene group, a pyrazinylene group, a pyrimidinylene group, a pyridazinylene group, an isoindolylene group, an indolylene group, an indazolylene group, a purinylene group, a quinolinylene group, an isoquinolinylene group, a benzoquinolinylene group, a phthalazinylene group, a naphthyridinylene group, a quinoxalinylene group, a quinazolinylene group, a cinnolinylene group, a carbazolylene group, a phenanthridinylene group, an acridinylene group, a phenanthrolinylene group, a phenazinylene group, a benzoimidazolylene group, a benzofuranylene group, a benzothiophenylene group, a benzoxazolylene group, an isobenzothiazolylene group, a benzoxazolylene group, a tetrazolylene group, an oxadiazolylene group, a triazinylene group, a dibenzofuranylene group, a dibenzothiophenylene group, a dibenzosilolylene group, a benzocarbazolylene group, a dibenzocarbazolylene group, a thiadiazolylene group, an imidazopyridinylene group, and an imidazopyrimidinylene group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a cyclopentyl group, a cyclohexyl group, a cycloheptyl group, a cyclopentenyl group, a cyclohexenyl group, a phenyl group, a biphenyl group, a terphenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an azulenyl group, a heptalenyl group, an indacenyl group, an acenaphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzo-

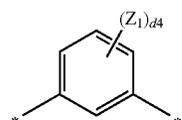
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fluorenyl group, a phenalenyl group, a phenanthrenyl group, an anthracenyl group, a fluoranthenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a naphthacenyl group, a picenyl group, a perylenyl group, a pentaphenyl group, a hexacenyl group, a pentacenyl group, a rubicenyl group, a coronenyl group, an ovalenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, an indolyl group, an indazolyl group, a purinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a phthalazinyl group, a naphthyridinyl group, a quinoxalinyl group, a quinazolinyl group, a cinnolinyl group, a carbazolyl group, a phenanthridinyl group, an acridinyl group, a phenanthrolinyl group, a phenazinyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolylene group, a benzocarbazolyl group, a dibenzocarbazolyl group, a thiadiazolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group.

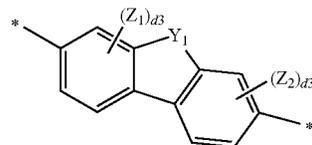
3. The organic light-emitting device of claim 1, wherein L₁ is selected from groups represented by Formula 3-1 to Formula 3-42:



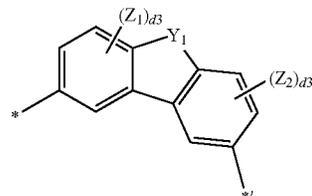
Formula 3-1



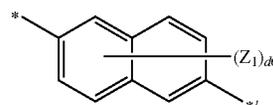
Formula 3-2



Formula 3-3

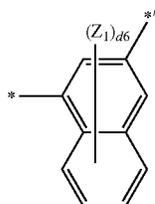
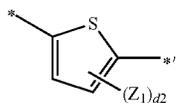
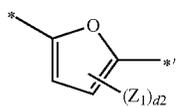
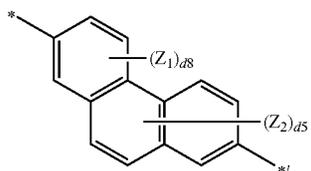
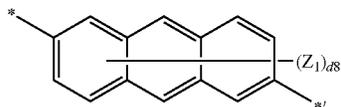
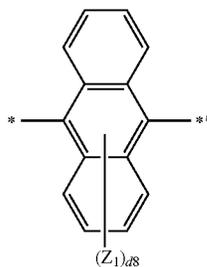
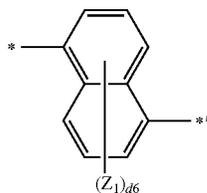
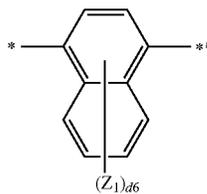


Formula 3-4



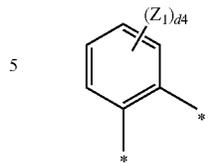
Formula 3-5

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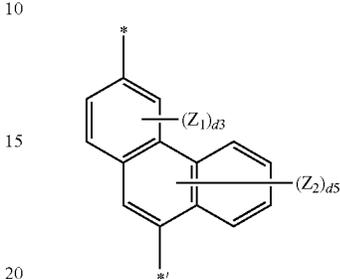


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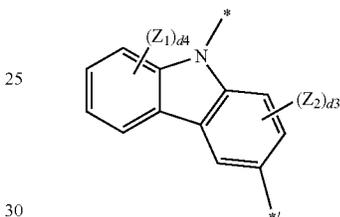
Formula 3-6



Formula 3-7

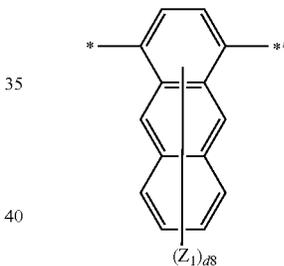


Formula 3-8

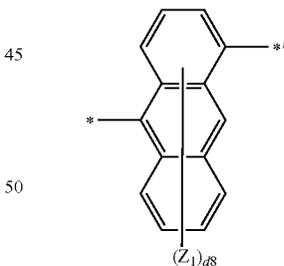


Formula 3-9

Formula 3-10



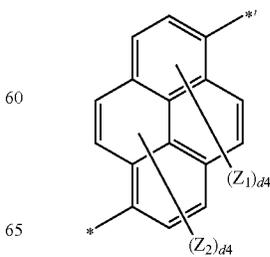
Formula 3-11



Formula 3-12

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Formula 3-13



Formula 3-14

Formula 3-15

Formula 3-16

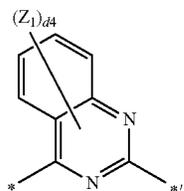
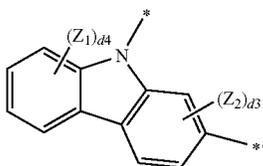
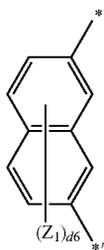
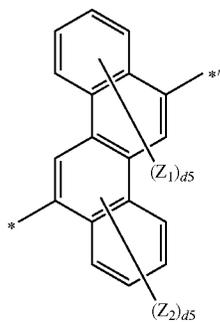
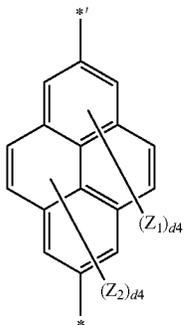
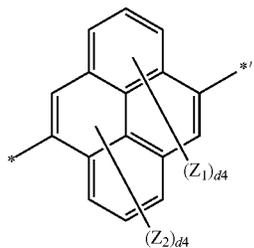
Formula 3-17

Formula 3-18

Formula 3-19

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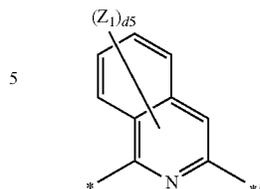
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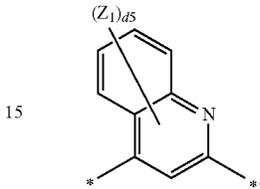
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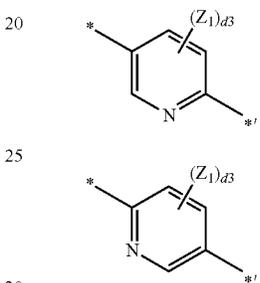
Formula 3-20



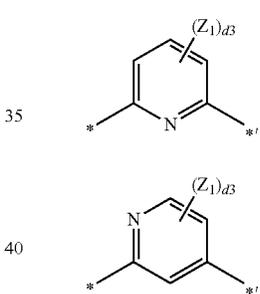
Formula 3-21



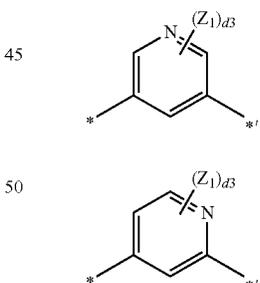
Formula 3-22



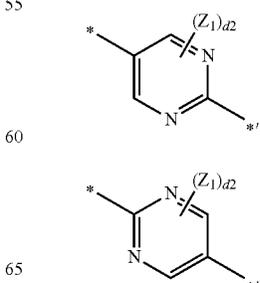
Formula 3-23



Formula 3-24



Formula 3-25



Formula 3-26

Formula 3-27

Formula 3-28

Formula 3-29

Formula 3-30

Formula 3-31

Formula 3-32

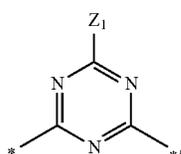
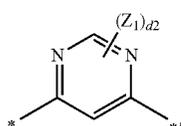
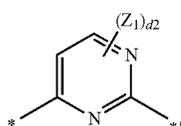
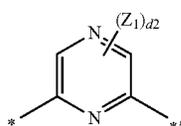
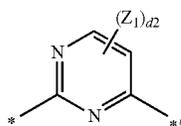
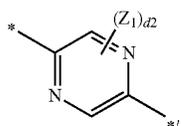
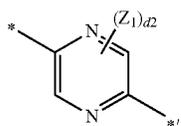
Formula 3-33

Formula 3-34

Formula 3-35

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wherein, in Formula 3-1 to Formula 3-42, Y_1 is selected from O, S, $C(Z_3)(Z_4)$, $N(Z_5)$, and $Si(Z_6)(Z_7)$;

Z_1 to Z_7 are each independently selected from hydrogen, deuterium, $-F$, $-Cl$, $-Br$, $-I$, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{20} alkyl group, a C_1 - C_{20} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a triazinyl group, and $-Si(Q_{31})(Q_{32})(Q_{33})$,

wherein Q_{31} to Q_{33} are each independently selected from a C_1 - C_{10} alkyl group, a C_1 - C_{10} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, and a naphthyl group;

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d2 is 1 or 2;

Formula 3-36

d3 is an integer selected from 1 to 3;

d4 is an integer selected from 1 to 4;

5

d5 is an integer selected from 1 to 5;

d6 is an integer selected from 1 to 6;

Formula 3-37

d8 is an integer selected from 1 to 8; and

10

* and *' each indicate a binding site to a neighboring atom.

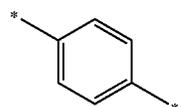
Formula 3-38

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4. The organic light-emitting device of claim 1, wherein Li is selected from groups represented by Formula 4-1 to Formula 4-29:

Formula 3-39

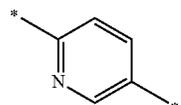
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Formula 4-1

Formula 3-40

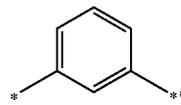
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Formula 4-2

Formula 3-41

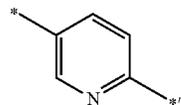
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Formula 4-3

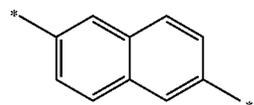
Formula 3-42

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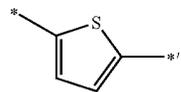
Formula 4-4

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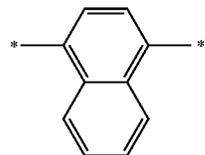
Formula 4-5

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Formula 4-6

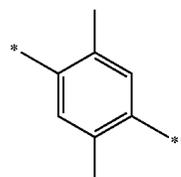
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Formula 4-7

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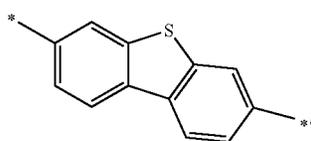
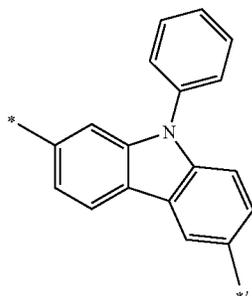
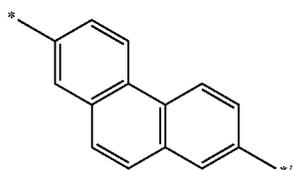
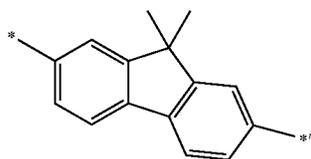
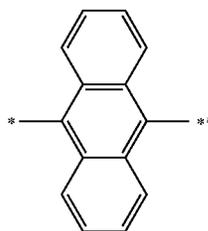
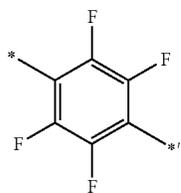
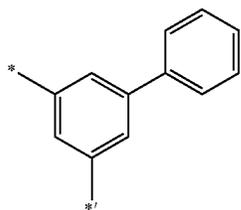


Formula 4-8

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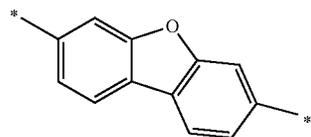


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Formula 4-9

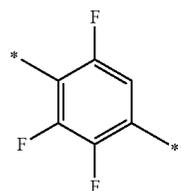
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Formula 4-16

Formula 4-10

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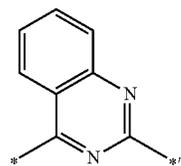


Formula 4-17

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Formula 4-11

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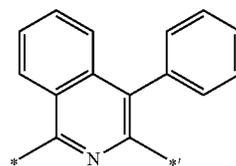


Formula 4-18

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Formula 4-12

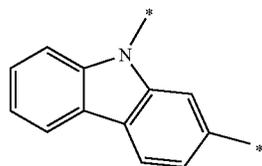
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Formula 4-19

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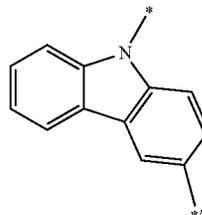
Formula 4-13



Formula 4-20

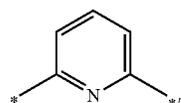
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Formula 4-14



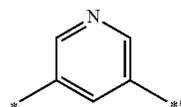
Formula 4-21

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Formula 4-22

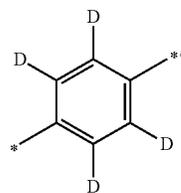
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Formula 4-23

Formula 4-15

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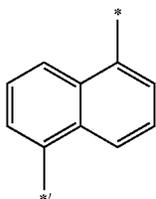
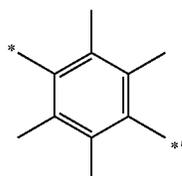
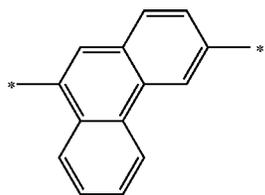
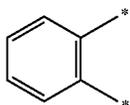
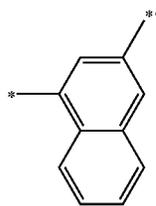


Formula 4-24

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wherein, in Formulae 4-1 to 4-29, * and *' each indicate a binding site to a neighboring atom.

5. The organic light-emitting device of claim 1, wherein L_1 is selected from the group consisting of:

a phenylene group, a naphthylene group, a fluorenylene group, a phenanthrenylene group, an anthracenylylene group, a triphenylenylene group, and a pyrenylene group, and

a phenylene group, a naphthylene group, a fluorenylene group, a phenanthrenylene group, an anthracenylylene group, a triphenylenylene group, and a pyrenylene group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an imino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{20} alkyl group, a C_1 - C_{20} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, and a naphthyl group.

6. The organic light-emitting device of claim 1, wherein a_1 is 0 or 1.

7. The organic light-emitting device of claim 1, wherein R_1 to R_6 and R_8 to R_{10} are each independently selected from the group consisting of:

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hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{20} alkyl group, and a C_1 - C_{20} alkoxy group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{10} alkyl group, a C_1 - C_{10} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group.

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group, an imidazopyridinyl group, an imidazopyrimidinyl group, and $-\text{Si}(\text{Q}_{31})(\text{Q}_{32})(\text{Q}_{33})$, and $-\text{Si}(\text{Q}_1)(\text{Q}_2)(\text{Q}_3)$,

wherein Q_1 to Q_3 and Q_{31} to Q_{33} are each independently selected from a C_1 - C_{10} alkyl group, a C_1 - C_{10} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, and a naphthyl group.

8. The organic light-emitting device of claim 1, wherein R_1 to R_6 and R_8 to R_{10} are each independently selected from the group consisting of:

hydrogen, deuterium, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, a hydroxyl group, a cyano group, a nitro group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{10} alkyl group, and a C_1 - C_{10} alkoxy group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a pyridinyl group, a pyrimidinyl group, and a triazinyl group,

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a pyridinyl group, a pyrimidinyl group, and a triazinyl group, each substituted with at least one selected from deuterium, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{10} alkyl group, a C_1 - C_{10} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a pyridinyl group, a pyrimidinyl group, a triazinyl group, and $-\text{Si}(\text{Q}_{31})(\text{Q}_{32})(\text{Q}_{33})$, and

$-\text{Si}(\text{Q}_1)(\text{Q}_2)(\text{Q}_3)$, wherein Q_1 to Q_3 and Q_{31} to Q_{33} are each independently selected from a C_1 - C_{10} alkyl group, a C_1 - C_{10} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, and a naphthyl group.

9. The organic light-emitting device of claim 1, wherein R_{11} is selected from the group consisting of:

a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiofenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group, and

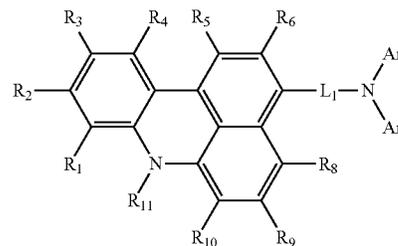
a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a

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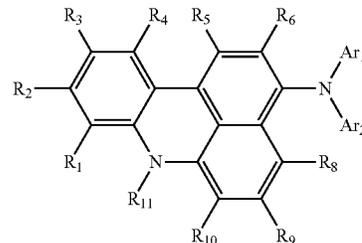
quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group, each substituted with at least one selected from deuterium, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{10} alkyl group, a C_1 - C_{10} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, an imidazopyridinyl group, an imidazopyrimidinyl group, and $-\text{Si}(\text{Q}_{31})(\text{Q}_{32})(\text{Q}_{33})$.

10. The organic light-emitting device of claim 1, wherein the condensed cyclic compound is represented by one selected from Formulae 1-2(A) and 1-2(B):

Formula 1-2(A)



Formula 1-2(B)



11. The organic light-emitting device of claim 10, wherein R_1 to R_6 and R_8 to R_{10} are each independently hydrogen; L_1 is selected from a substituted or unsubstituted phenylene group, a substituted or unsubstituted naphthyl-

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ene group, a substituted or unsubstituted fluorenylene group, a substituted or unsubstituted phenanthrenylene group, a substituted or unsubstituted anthracenylylene group, a substituted or unsubstituted triphenylylene group, and a substituted or unsubstituted pyrenylene group;

R_{11} is selected from a substituted or unsubstituted phenyl group, a substituted or unsubstituted biphenyl group, a substituted or unsubstituted terphenyl group, a substituted or unsubstituted naphthyl group, a substituted or unsubstituted fluorenyl group, a substituted or unsubstituted phenanthrenyl group, a substituted or unsubstituted anthracenyl group, a substituted or unsubstituted triphenylenyl group, a substituted or unsubstituted pyrenyl group, a substituted or unsubstituted fluorenyl group, a substituted or unsubstituted spiro-bifluorenyl group, a substituted or unsubstituted carbazolyl group, a substituted or unsubstituted dibenzofuranyl group, a substituted or unsubstituted dibenzothiophenyl group, and a substituted or unsubstituted dibenzosilolyl group; and

at least one substituent of the substituted phenylene group, substituted naphthylene group, substituted fluorenylene group, substituted phenanthrenylene group, substituted anthracenylylene group, substituted triphenylylene group, substituted pyrenylene group, substituted phenyl group, substituted biphenyl group, substituted terphenyl group, substituted naphthyl group, substituted fluorenyl group, substituted phenanthrenyl group, substituted anthracenyl group, substituted triphenylenyl group, substituted pyrenyl group, substituted fluorenyl group, substituted spiro-bifluorenyl group, substituted carbazolyl group, substituted dibenzofuranyl group, substituted dibenzothiophenyl group, and substituted dibenzosilolyl group is selected from the group consisting of:

hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{20} alkyl group, and a C_1 - C_{20} alkoxy group,

a C_1 - C_{20} alkyl group, and a C_1 - C_{20} alkoxy group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, and a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a carbazolyl group, a benzocarbazolyl group, and a dibenzocarbazolyl group,

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a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a pyridinyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a carbazolyl group, a benzocarbazolyl group, and a dibenzocarbazolyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{20} alkyl group, a C_1 - C_{20} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, and a pyridinyl group, and

—Si(Q_{31})(Q_{32})(Q_{33}),

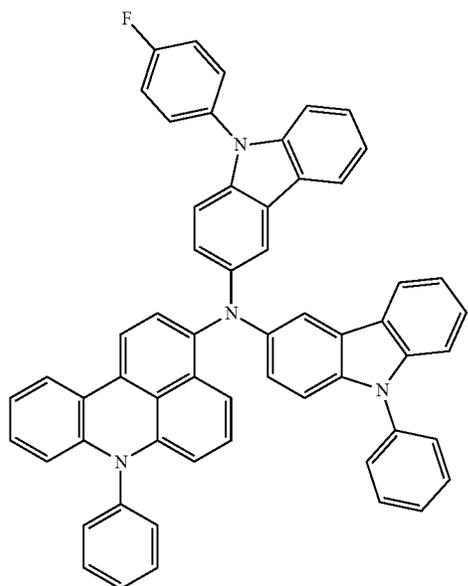
wherein Q_{31} to Q_{33} are each independently selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{20} alkyl group, a C_1 - C_{20} alkoxy group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, and a carbazolyl group.

12. An organic light-emitting device comprising:

- a first electrode;
- a second electrode facing the first electrode; and
- an organic layer between the first electrode and the second electrode, the organic layer comprising:
 - i) an emission layer,
 - ii) a hole transport region between the first electrode and the emission layer, the hole transport region comprising at least one selected from a hole injection layer, a hole transport layer, a buffer layer, and an electron blocking layer, and
 - iii) an electron transport region between the emission layer and the second electrode, the electron transport region comprising at least one selected from a hole blocking layer, an electron transport layer, and an electron injection layer,

wherein the emission layer is to emit blue fluorescence, wherein the hole transport region comprises a condensed cyclic compound selected from Compounds 57, 66, 81, and 90:

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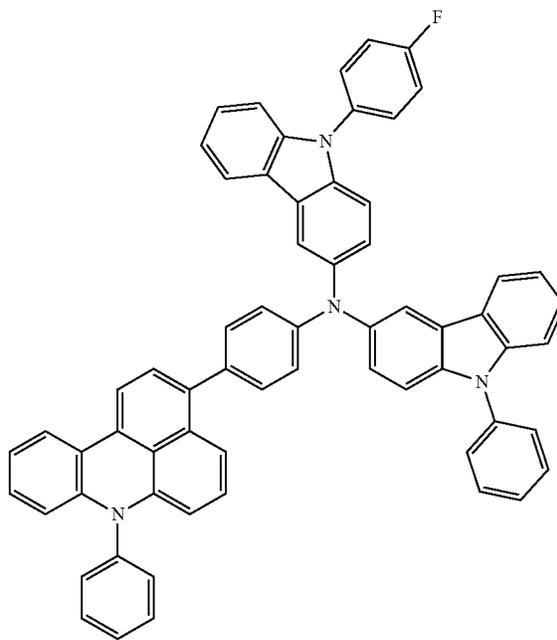
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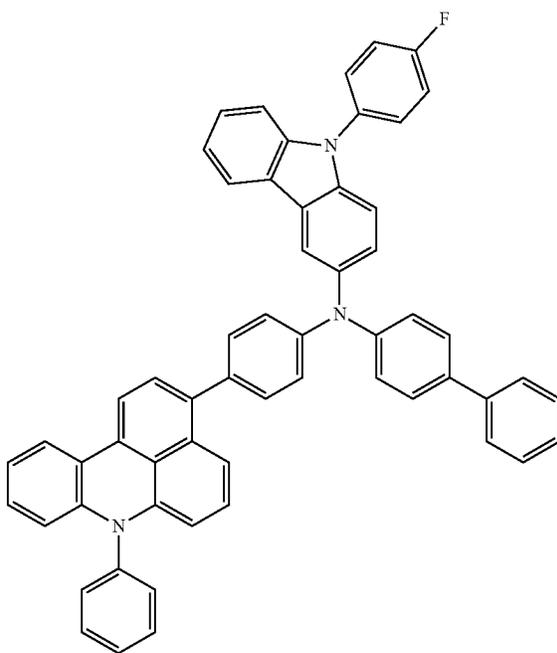
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13. The organic light-emitting device of claim 1, wherein the hole transport region comprises a charge-generating material.

* * * * *